



2017 EACTS/EACTA Guidelines on patient blood management for adult cardiac surgery

Boer, Christa; Meesters, Michael I; Milojevic, Milan; Benedetto, Umberto; Bolliger, Daniel; von Heymann, Christian; Jeppsson, Anders; Koster, Andreas; Osnabrugge, Ruben L; Ranucci, Marco; Ravn, Hanne Berg; Vonk, Alexander B A; Wahba, Alexander; Pagano, Domenico; Task Force on Patient Blood Management for Adult Cardiac Surgery of the European Association for Cardio-Thoracic Surgery (EACTS) and the European Association of Cardiothoracic Anaesthesiology (EACTA)

Published in:
Journal of Cardiothoracic and Vascular Anesthesia

DOI:
[10.1053/j.jvca.2017.06.026](https://doi.org/10.1053/j.jvca.2017.06.026)

Publication date:
2018

Document version
Publisher's PDF, also known as Version of record

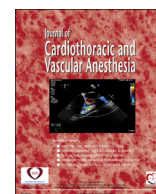
Document license:
[CC BY](#)

Citation for published version (APA):
Boer, C., Meesters, M. I., Milojevic, M., Benedetto, U., Bolliger, D., von Heymann, C., Jeppsson, A., Koster, A., Osnabrugge, R. L., Ranucci, M., Ravn, H. B., Vonk, A. B. A., Wahba, A., Pagano, D., & Task Force on Patient Blood Management for Adult Cardiac Surgery of the European Association for Cardio-Thoracic Surgery (EACTS) and the European Association of Cardiothoracic Anaesthesiology (EACTA) (2018). 2017 EACTS/EACTA Guidelines on patient blood management for adult cardiac surgery. *Journal of Cardiothoracic and Vascular Anesthesia*, 32(1), 88-120. <https://doi.org/10.1053/j.jvca.2017.06.026>



Contents lists available at ScienceDirect

ScienceDirect

journal homepage: www.jcvaonline.com

2017 EACTS/EACTA Guidelines on patient blood management for adult cardiac surgery



The Task Force on Patient Blood Management for Adult Cardiac Surgery
of the European Association for Cardio-Thoracic Surgery (EACTS) and the
European Association of Cardiothoracic Anaesthesiology (EACTA)

Christa Boer (EACTA Chairperson)^{*,1,2}, Michael I. Meesters^{1,2}, Milan Milojevic¹,
Umberto Benedetto¹, Daniel Bolliger^{1,2}, Christian von Heymann^{1,2}, Anders Jeppsson¹,
Andreas Koster^{1,2}, Ruben L. Osnabrugge¹, Marco Ranucci^{1,2}, Hanne Berg Ravn^{1,2},
Alexander B.A. Vonk¹, Alexander Wahba¹, Domenico Pagano (EACTS Chairperson)^{*,1}

Key Words: Guidelines; Cardiac surgery; Haemostasis; Blood transfusion; Bleeding; Anticoagulation; Coagulation factors

Document Reviewers: Moritz W.V. Wyler von Ballmoos (USA), Mate Petricevic (Croatia), Arie Pieter Kappetein (Netherlands), Miguel Sousa-Uva (Portugal), Georg Trummer (Germany), Peter M. Rosseel (Netherlands)², Michael Sander (Germany)², Pascal Colson (France)², Adrian Bauer (Germany)³

Table of Contents

Abbreviations and Acronyms	89
1. Preamble	90
2. Introduction	90
3. Methods	90
4. Preoperative Management.	91

Disclaimer 2017: The EACTS/EACTA Guidelines represent the views of the EACTS and of the EACTA and were produced after careful consideration of the scientific and medical knowledge and the evidence available at the time of their dating.

The EACTS and the EACTA are not responsible in the event of any contradiction, discrepancy and/or ambiguity between the EACTS/EACTA Guidelines and any other official recommendations or guidelines issued by the relevant public health authorities, in particular in relation to good use of healthcare or therapeutic strategies. Health professionals are encouraged to take the EACTS/EACTA Guidelines fully into account when exercising their clinical judgement as well as in the determination and the implementation of preventive, diagnostic or therapeutic medical strategies; however, the EACTS/EACTA Guidelines do not, in any way whatsoever, override the individual responsibility of health professionals to make appropriate and accurate decisions in consideration of each patient's health condition and, where appropriate and/or necessary, in consultation with that patient and the patient's care provider. Nor do the EACTS/EACTA Guidelines exempt health professionals from giving full and careful consideration to the relevant official, updated recommendations or guidelines issued by the competent public health authorities, in order to manage each patient's case in light of the scientifically accepted data pursuant to their respective ethical and professional obligations. It is also the health professional's responsibility to verify the applicable rules and regulations relating to drugs and medical devices at the time of prescription.

The article has been co-published with permission in the European Journal of Cardio-Thoracic Surgery and in the Journal of Cardiothoracic and Vascular Anesthesia. All rights reserved in respect of the European Journal of Cardio-Thoracic Surgery, © European Association for Cardio-Thoracic Surgery 2017 and in respect of the Journal of Cardiothoracic and Vascular Anesthesia © European Association of Cardiothoracic Anaesthesiology 2017. The articles are identical except for minor stylistic and spelling difference in keeping with each journal's style. Either citation can be used when citing this article.

*Address reprint requests to: Christa Boer, Department of Anesthesiology, VU University Medical Center, Amsterdam, Netherlands. Tel.: +31-20-4443830; Domenico Pagano, Department of Cardiothoracic Surgery, University Hospital Birmingham, Birmingham, UK. Tel.: +44-121-6272850.

E-mail addresses: c.boer@vumc.nl (C. Boer), domenico.pagano@uhb.nhs.uk (D. Pagano).

¹Authors/Task Force Members

²Representing the EACTA

³Representing the European Board of Cardiovascular Perfusion (EBCP)

4.1.	Laboratory and point-of-care tests to predict perioperative bleeding	91	7.	Anticoagulation Management During Extracorporeal Life Support.	111
4.2.	Management of preoperative anticoagulant and antiplatelet drugs	92	8.	Final Remarks	111
4.2.1.	Acetylsalicylic acid	93		References	112
4.2.2.	Dual antiplatelet therapy	93	Abbreviations and Acronyms		
4.2.3.	Glycoprotein IIb/IIIa inhibitors	94	ACS	Acute coronary syndrome	
4.2.4.	Low-molecular-weight heparin	95	ACT	Activated clotting time	
4.2.5.	Vitamin K antagonists	95	AKI	Acute kidney injury	
4.2.6.	Direct oral anticoagulant	96	ANH	Acute normovolaemic haemodilution	
4.3.	Preoperative anaemia	96	aPTT	Activated partial thromboplastin time	
4.3.1.	Implications for preoperative anaemia	96	ASA	Acetylsalicylic acid	
4.3.2.	Iron supplementation	96	AT	Antithrombin	
4.3.3.	Erythropoietin	97	CABG	Coronary artery bypass grafting	
4.3.4.	Blood transfusion to treat preoperative anaemia	97	CI	Confidence interval	
5.	Intraoperative Management.	97	CKD	Chronic kidney disease	
5.1.	Surgical techniques.	97	CPB	Cardiopulmonary bypass	
5.1.1.	Off-pump surgery	98	DAPT	Dual antiplatelet therapy	
5.1.2.	Minimally invasive extracorporeal circulation.	98	DDAVP	Desmopressin	
5.1.3.	Minimally invasive surgery	98	DOAC	Direct oral anticoagulant	
5.1.4.	Topical haemostatics.	99	EACA	ϵ -Aminocaproic acid	
5.2.	Cardiopulmonary bypass	99	EACTA	European Association of Cardiothoracic Anaesthesiology	
5.2.1.	Closed versus open cardiopulmonary bypass	99	EACTS	European Association for Cardio-Thoracic Surgery	
5.2.2.	Biocompatible coating	100	ECC	Extracorporeal circulation	
5.2.3.	Cell salvage	100	ECLS	Extracorporeal life support	
5.2.4.	Ultrafiltration.	101	ECMO	Extracorporeal membrane oxygenation	
5.2.5.	Retrograde and antegrade autologous priming	101	EPO	Erythropoietin	
5.2.6.	Coagulation-friendly environment.	101	FFP	Fresh-frozen plasma	
5.3.	Intraoperative anticoagulation.	101	FXIII	Factor XIII	
5.3.1.	Heparin and anticoagulation monitoring.	101	GPIIb/IIIa	Glycoprotein	
5.3.2.	Protamine	102	Hb	Haemoglobin	
5.3.3.	Antithrombin.	102	HES	Hydroxyethyl starches	
5.3.4.	Heparin-induced thrombocytopenia anticoagulation management	103	HIT	Heparin-induced thrombocytopenia	
5.4.	Intravascular volume.	103	HR	Hazard ratio	
5.4.1.	Goal-directed haemodynamic therapy	103	INR	International normalized ratio	
5.4.2.	Use of crystalloids and colloids	104	LMWH	Low-molecular-weight heparin	
5.4.3.	Haemodilution and cardioplegia	104	MI	Myocardial infarction	
5.4.4.	Predonation of blood	104	MiECC	Minimally invasive extracorporeal circulation circuit	
5.4.5.	Acute normovolaemic haemodilution.	105	OR	Odds ratio	
6.	Coagulation and Transfusion.	105	PBM	Patient blood management	
6.1.	Procoagulant interventions.	105	PCC	Prothrombin complex concentrate	
6.1.1.	Antifibrinolytics.	105	PLTC	Platelet concentrate	
6.1.2.	Fresh-frozen plasma	106	PMEA	Poly2-methoxyethylacrylate	
6.1.3.	Factor XIII	106	POC	Point-of-care	
6.1.4.	Fibrinogen concentrates.	106	PRBC	Packed red blood cells	
6.1.5.	Prothrombin complex concentrate.	107	RAP	Retrograde autologous priming	
6.1.6.	Desmopressin	107	RCT	Randomized controlled trial	
6.1.7.	Recombinant factor VIIa.	108	rFVIIa	Recombinant activated factor seven	
6.2.	Transfusion strategies	108	RR	Risk ratio	
6.2.1.	Quality of blood products	108	SD	Solvent–detergent	
6.2.2.	Algorithm-guided therapy of perioperative bleeding	109			
6.2.3.	Transfusion triggers for packed red blood cells and platelet concentrate	110			

TEG	Thromboelastography
TEM	Thromboelastometry
TRALI	Transfusion-related acute lung injury
TRIM	Transfusion-related immune modulation
TXA	Tranexamic acid
UFH	Unfractionated heparin
VKA	Vitamin K antagonist

1. Preamble

Cardiac surgery is associated with perioperative blood loss and a high risk of allogeneic blood transfusion. Patient blood management (PBM) in cardiac surgery contributes to the maintenance of perioperative haemostasis and the minimization of bleeding, which reduce blood transfusion requirements. PBM in cardiac surgery comprises an interaction between the cardiothoracic surgeon, the anaesthesiologist and the clinical perfusionist. The impact of cardiopulmonary bypass distinguishes this discipline from other surgical specialities.

In a joint effort, the European Association for Cardio-Thoracic Surgery (EACTS) and the European Association of Cardiothoracic Anaesthesiology (EACTA) provide evidence-based recommendations for PBM in adult-acquired cardiac surgery. Literature searches were based on the Population, Intervention, Comparison, Outcome and Time (PICOT) method using standardized Medical Subject Headings (MeSH) terms from the National Library of Medicine, PubMed and Embase database lists of search terms. The PICOT study end points included bleeding, transfusions and reoperations for bleeding. The guideline was reviewed by an external review method and endorsed by the EACTS and the EACTA in collaboration with the editors of the *European Journal of Cardio-Thoracic Surgery* and the *Journal of Cardiothoracic and Vascular Anesthesia*.

This guideline provides practical recommendations for all clinicians working in the field of PBM in cardiac surgery, with emphasis on preoperative patient optimization and risk reduction, intraoperative maintenance of haemostasis and postoperative treatment for bleeding complications.

2. Introduction

Cardiac surgery is known to be associated with a high risk of perioperative blood loss and allogeneic blood transfusions due to the invasiveness of the procedures, the need for high-dose anticoagulation and the exposure to cardiopulmonary bypass (CPB). Both high blood product transfusion requirements and reoperation for bleeding have been associated with adverse clinical outcomes.^{1,2} The implementation of a multidisciplinary patient blood management (PBM) programme, the goal of which is meticulous surgical and perioperative haemostasis and minimization of blood loss, may, therefore, contribute to a reduction in transfusion requirements, a decrease in health care costs and an improvement in patient outcomes.

Even the transfusion of 1 or 2 units of packed red blood cells (PRBCs) has been associated with a dramatic increase in morbidity, mortality and costs in patients undergoing coronary artery bypass grafting (CABG).³ However, it remains unclear whether these complications are independent predictors of outcome or more surrogate markers of surgical complexity and complications. In contrast, re-exploration for bleeding and tamponade appears to be a strong risk factor for an increase in immediate postoperative mortality and morbidity. There is, however, no general consensus on when exploration for postoperative bleeding is indicated, and surgical practice varies considerably in this regard.⁴ The first step towards creating algorithms to decrease blood loss and transfusions is to identify patients at high risk of bleeding, transfusion requirements and reoperation for bleeding. Also, there is little consensus on what constitutes a massive blood transfusion and what thresholds might be associated with adverse outcomes.^{5,6}

Several factors have been found to be associated with an increased risk of bleeding, transfusion and reoperation,^{7–9} including advanced age, preoperative dual antiplatelet therapy (DAPT) use, poor platelet function, preoperative anaemia, small body surface area, the female gender, non-elective surgery, nonisolated surgery, non-CABG surgery and redo surgery. In an attempt to facilitate the preoperative prediction of the need for transfusion, several scoring methods have been developed.^{7,10–15} However, most of these risk models are of uncertain relevance, because they have not been externally validated. Risk prediction algorithms are, therefore, mostly used to stratify outcomes and allow risk-adjusted benchmarking in the context of quality and outcome monitoring.

PBM in cardiac surgery comprises an interaction between the surgeon, the anaesthesiologist and the clinical perfusionist, and the use of CPB distinguishes this discipline from other surgical specialities. Despite the availability of recent guidelines on PBM,^{9,16,17} there are currently no PBM guidelines available that specifically address the unique context of cardiac surgery and that are endorsed by the European societies. In a joint effort, the European Association for Cardio-Thoracic Surgery (EACTS) and the European Association of Cardiothoracic Anaesthesiology (EACTA) collaborated on a common guideline with evidence-based recommendations for PBM in adult-acquired cardiac surgery conditions.

3. Methods

As an expression of their declared and consolidated interest in medical care and scientific research in the field of PBM in cardiac surgery, the EACTS Council and the Board of Directors of EACTA selected a task force of experts, including a clinical methodologist. After the scope of the clinical guideline had been agreed upon by the task force members, a preliminary review of previous guidelines was performed. The scope and table of contents were established by co-chairs, and topics were allocated to the writing group. A systematic review of the published evidence in the field of blood conservation during adult-acquired cardiac surgery was

performed based on the Population, Intervention, Comparison, Outcome and Time (PICOT) questions. Task force members undertook an evidence review; they were assisted by 2 research fellows with an MD/PhD in anaesthesiology and an MSc in clinical epidemiology, respectively. After study selection and quality assessment, tables of recommendations were drafted based on the synthesis of the best medical evidence. The experts conducted a review of the available evidence and followed the Methodology Manual for EACTS clinical guidelines.¹⁸ All chapters were written through a collaboration between an EACTS member and an EACTA member. Agreement was reached through conference calls and face-to-face meetings, without excluding members with a conflict of interest. When no agreement could be obtained, a consensus was reached following a modified Delphi process.¹⁹ Task force members who were unable to attend a face-to-face meeting voted by email. Three Delphi rounds were required to reach a consensus, and a final decision was made through a series of teleconference calls. Chapters were written by task force members who did not have a disclosure for the specific topic.

The literature was restricted to the years 2001–2017 to focus on contemporary evidence. The guideline only focused on adult-acquired cardiac surgery and did not include studies in the areas of transplantation, trauma, circulatory arrest or long-term circulatory support. Outcomes were defined as bleeding volume or chest drain output in the first 24 h following surgery, transfusion requirements and reoperation for bleeding at any time point during hospitalization. The definitions of bleeding, time frames of measurement and thresholds for transfusions varied significantly between studies. Wherever a distinction in time frame beyond 24 h was made, it is emphasized in the text.

The medical evidence was critically appraised for quality, including internal validity, external validity for the population of interest and publication bias. Randomized controlled trials (RCTs) and meta-analyses of RCTs were considered as the highest level of evidence, followed by other study designs. When a meta-analysis was the basis for a recommendation, the quality was assessed by a group of specialists from the EACTS/EACTA. In the absence of published evidence, expert consensus statements were made to cover specific issues that are essential to daily practice. The level of evidence and the strength of the recommendations were weighed and graded according to pre-defined scales, as outlined in Table 1 and 2.

Table 1
Classes of recommendations

Classes of recommendations	Definition	Suggested wording to use
Class I	Evidence and/or general agreement that a given treatment or procedure is beneficial, useful and effective.	Is recommended/is indicated
Class II	Conflicting evidence and/or a divergence of opinion about the usefulness/efficacy of the given treatment or procedure.	
Class IIa	Weight of evidence/opinion is in favour of usefulness/efficacy.	Should be considered
Class IIb	Usefulness/efficacy is less well established by evidence/opinion.	May be considered
Class III	Evidence/general agreement that the given treatment/procedure is not useful/effective and may sometimes be harmful.	Is not recommended

Classes of recommendations based on the Methodology Manual for European Association for Cardio-Thoracic Surgery clinical guidelines.¹⁸

Table 2
Levels of evidence

Level of evidence A	Data derived from multiple randomized clinical trials or meta-analyses.
Level of evidence B	Data derived from a single randomized clinical trial or large non-randomized studies.
Level of evidence C	The consensus of expert opinion and/or small studies, retrospective studies, registries.

Levels of evidence based on the Methodology Manual for European Association for Cardio-Thoracic Surgery clinical guidelines.¹⁸

4. Preoperative Management

Inadequate patient optimization before surgery increases the risk of intraoperative anaemia and bleeding. Preoperative measures include management of antithrombotic medication, optimization of haemoglobin (Hb) levels in the context of patient comorbidities and body surface dimensions and the assessment and weighing of patient haemostatic risk factors, including the presence of congenital disorders.

4.1. Laboratory and Point-of-Care Tests to Predict Perioperative Bleeding

Recommendations	Class ^a	Level ^b	Ref ^c
Preoperative fibrinogen levels may be considered to identify patients at high risk of bleeding.	IIb	C	
Routine use of viscoelastic and platelet function testing is not recommended to predict bleeding in patients without antithrombotic treatment.	III	C	
Platelet function testing may be considered to guide the decision on the timing of cardiac surgery in patients who have recently received P2Y12 inhibitors or who have ongoing DAPT.	IIb	B	20–22

^aClass of recommendation.

^bLevel of evidence.

^cReferences.

DAPT: dual antiplatelet therapy.

Background

Preoperative assessment of haemostatic parameters by standard laboratory testing, viscoelastic testing and/or platelet function tests has been proposed to predict perioperative bleeding in cardiac surgery.

Description of the Evidence

Standard Laboratory Testing. The use of routine preoperative screening has been much debated regarding its ability to identify high-risk patients for postoperative bleeding and transfusion requirements. There is no association between preoperative prothrombin time or the activated partial thromboplastin time (aPTT) and perioperative blood loss or transfusion requirements.^{23–25} So far, the most commonly identified risk factor for postoperative bleeding is a lowfibrinogen level.^{23,26–28} However, despite its association with bleeding, the positive predictive value of a low-fibrinogen level remains poor (positive predictive value < 20%).²⁶ A low-platelet count < 100 × 10⁹/l has been associated with increased risk of transfusion,¹² and patients with the highest postoperative blood loss volumes show the lowest platelet counts.²⁴ Finally, it has been shown that patients with the highest postoperative blood loss volumes show the lowest thrombin generation rates,^{24,29} but this test is mainly used for research purposes and is not routinely used in everyday practice.

Preoperative fibrinogen levels and thrombin generation assays may be considered to identify patients at high risk of bleeding, e.g. during aortic or emergency surgery, although thrombin generation assays are not yet validated for routine clinical application.

Viscoelastic Testing and Platelet Function Tests. Preoperative assessment of haemostatic parameters using viscoelastic tests, such as rotational thromboelastometry (TEM) or thromboelastography (TEG), has been found to have a limited association with the risk of postoperative bleeding.^{29–31} Preoperative platelet function testing has not been demonstrated to be associated with an increased risk of bleeding complications in patients without ongoing or recently stopped DAPT. In contrast, several studies have shown a significant association between impaired platelet function and bleeding complications in patients on DAPT. This finding indicates that platelet function testing may be used to guide the timing of surgery in this group of patients.^{21,22,32,33}

4.2. Management of Preoperative Anticoagulant and Antiplatelet Drugs

Recommendations	Class ^a	Level ^b	Ref ^c
In patients undergoing CABG, ASA should be continued throughout the preoperative period.	IIa	C	
In patients at high risk of bleeding ^d or refusing blood transfusions and undergoing non-coronary cardiac surgery, stopping ASA should be considered at least 5 days preoperatively.	IIa	C	

It is recommended that ASA be re(started) as soon as there is no concern over bleeding (within 24 h) after isolated CABG.	I	B	34,35
In patients taking DAPT who need to have non-emergent cardiac surgery, postponing surgery for at least 3 days after discontinuation of ticagrelor, 5 days after clopidogrel and 7 days after prasugrel should be considered.	IIa	B	36–38
It is recommended that GPIIb/IIIa inhibitors be discontinued at least 4 h before surgery.	I	C	
To reduce the risk of bleeding, preoperative bridging of oral anticoagulation with UFH/LMWH is only indicated in patients at high risk of thrombotic events. ^e	I	B	39
It is recommended that prophylactic LMWH be discontinued 12 h before surgery and fondaparinux 24 h before surgery. A longer interval may be necessary for patients with impaired renal function and/ or therapeutic doses.	I	B	40–42
It is recommended that OACs be bridged with UFH.	I	B	43,44
Bridging OACs with subcutaneous LMWH should be considered an alternative to bridging with UFH.	IIa	B	45,46
Elective cardiac surgery should be performed if the INR is < 1.5 in patients taking VKAs. When surgery cannot be postponed, coagulation factors should be used to reverse the effect.	IIa	C	
In patients having elective cardiac surgery, DOACs should be stopped at least 48 h before surgery. A longer interval may be necessary for patients with impaired renal function.	IIa	C	

^aClass of recommendation.

^bLevel of evidence.

^cReferences.

^dComplex and redo operation, severe renal insufficiency, haematological diseases and hereditary deficiencies in platelet function.

^eMechanical prosthetic heart valve, atrial fibrillation with rheumatic valvular disease, an acute thrombotic event within the previous 4 weeks and atrial fibrillation with a CHA₂DS₂-VASc score > 4.

ASA: acetylsalicylic acid; CABG: coronary artery bypass grafting; CHA₂DS₂-VASc: congestive heart failure, hypertension, age ≥ 75 (2 points), diabetes, prior stroke (2 points) – vascular disease, age 65–74, sex category (female); DAPT: dual antiplatelet therapy; DOAC: direct oral anticoagulant; GP: glycoprotein; INR: international normalized ratio; LMWH: low-molecular-weight heparin; OAC: oral anticoagulants; UFH: unfractionated heparin; VKA: vitamin K antagonist.

4.2.1. Acetylsalicylic Acid

Background. Acetylsalicylic acid (ASA) is one of the cornerstones for the treatment of acute and chronic cardiovascular disease. Primary and secondary prevention with ASA has been shown to reduce mortality, myocardial infarction (MI) and stroke⁴⁷ but to increase the risk of bleeding complications. Virtually, all patients requiring CABG—independent of whether they require emergent, urgent or elective surgery—are treated with ASA. Figure 1 provides an overview of the management of antiplatelet therapy in patients undergoing CABG surgery.

Description of the Evidence

Discontinuation before Surgery. A recent meta-analysis comparing preoperative ASA administration with distinct dosing regimens versus no treatment or placebo in patients having CABG included 13 RCTs ($n = 2399$ patients).⁴⁸ The meta-analysis showed that ASA reduced the risk of perioperative MI [odds ratio (OR) 0.56, 95% confidence interval (CI) 0.33–0.96] but not the risk of death (OR 1.16, 95% CI 0.42–3.22). Twelve-hour blood loss, PRBC transfusions and surgical re-exploration increased with ASA. The meta-analysis was limited by substantial heterogeneity but confirmed findings from a previous meta-analysis.⁴⁹ A large RCT compared the administration of ASA (100 mg) on the day of surgery versus placebo in patients having CABG.⁵⁰ The study showed no effect of treatment with ASA on 24-h bleeding (mean blood loss: 780 vs 740 ml; $P = 0.30$) or on the incidence of death or thrombotic complications (19.3% vs 20.4%; $P = 0.55$). Because patients were only eligible for inclusion if they were not using ASA or stopped ASA > 4 days before surgery, these findings are difficult to generalize to other settings.⁵⁰ Another RCT demonstrated that preoperative administration of ASA (300 mg) was associated with significantly more patients having post-surgical drainage losses > 1000 ml (OR 1.60,

95% CI 1.17–2.18) but with a lower rate of major cardiovascular events at the 53-month follow-up compared with placebo [hazard ratio (HR) 0.65, 95% CI 0.41–1.03].⁵¹ In a small RCT in patients with a platelet glycoprotein (GP) IIIa polymorphism, treatment with ASA (300 mg) increased the risk of postoperative bleeding (+25%).⁵²

If one considers all of the results from the different studies, continuation of ASA is associated with more blood loss but also reduces ischaemic events in patients having CABG. In patients who refuse blood transfusions, who are undergoing non-CABG surgery or who are at a high risk of re-exploration for bleeding (e.g. in cases of complex and redo operations, severe renal insufficiency, haematological disease and hereditary platelet function deficiencies), ASA should be stopped at least 5 days before surgery. In other patients, the prevention of thrombotic events outweighs the risk of postoperative bleeding events. Moreover, recent data suggest that the inhibiting effect of ASA on platelet aggregation is reversible by platelet transfusion,^{53,54} which also argues for the continuation of ASA.

Because current evidence supports the idea that ASA should be continued before surgery, the question of restarting applies only to a minority of patients. In a large prospective observational trial,³⁴ patients who restarted ASA within 48 h of CABG had a mortality rate of 1.3% compared to 4.0% in the group of patients who did not receive ASA during this period ($P < 0.001$).

In conclusion, there is sufficient evidence to recommend continuation of ASA before cardiac surgery, and, if discontinued, it should be given to all patients having CABG as soon as there is no concern over bleeding to prevent thromboembolic complications.

4.2.2. Dual Antiplatelet Therapy

Background. DAPT with ASA and a P2Y₁₂-receptor antagonist (clopidogrel, ticagrelor and prasugrel) reduces the

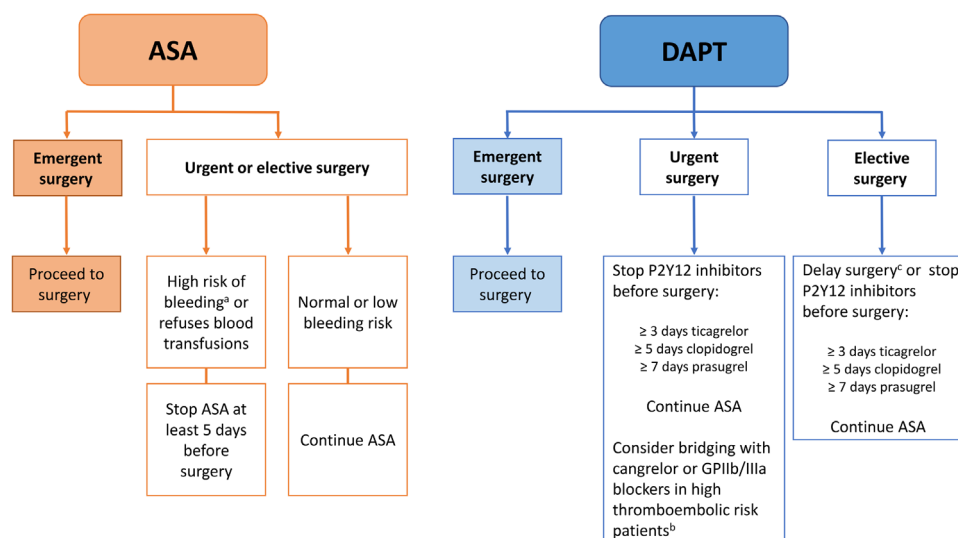


Figure 1. Management of antiplatelet therapy in patients having coronary artery bypass grafting surgery. ^aComplex and redo operations, severe renal insufficiency, haematological diseases and hereditary deficiencies in platelet function. ^bRecent stent implantation, recent thromboembolic event and alarming angiographic results. ^cUntil the recommended DAPT period is completed. ASA: acetylsalicylic acid; DAPT: dual antiplatelet therapy; GPIIb/IIIa: glycoprotein IIb/IIIa.

risk of thrombotic complications in patients with acute coronary syndrome (ACS) compared with ASA treatment only.^{55–57} The risk of thrombotic complications is further reduced if a second-generation P2Y₁₂ antagonist (ticagrelor/prasugrel) is used instead of clopidogrel,^{56,57} but the risk of both spontaneous and surgical bleeding complications increases with the newer antagonists.^{56–58} The duration of DAPT treatment varies depending on the indication (recent ACS and/or stent implantation, coronary endarterectomy or off-pump surgery), bleeding risks and concomitant medications.⁵⁹ Recently, cangrelor, a new reversible intravenous P2Y₁₂ inhibitor with an ultrashort half-life to offset the effect after discontinuation, was introduced.⁶⁰

Description of the Evidence

Discontinuation of P2Y₁₂ Inhibitors Before Surgery. Continuation of DAPT until surgery increases the risk of postoperative bleeding, transfusions and re-exploration for bleeding, as shown in RCTs,^{36,61,62} observational studies^{37,63} and meta-analyses.^{64,65} It is therefore recommended that P2Y₁₂ inhibitors be discontinued before elective surgery whenever feasible.^{66,67} Of note is the fact that, if P2Y₁₂-receptor inhibitors are discontinued, ASA therapy should be continued until surgery. Alternatively, elective non-cardiac and cardiac procedures may be postponed until the DAPT treatment period is completed. In urgent cases, the risk of thromboembolic episodes must be weighed against the risk of perioperative bleeding complications. In extremely high-risk patients (e.g. patients with recent stent implantation), bridging therapy with cangrelor or a GPIIb/IIIa blocker may be considered.^{66,67}

The safe discontinuation interval differs among distinct P2Y₁₂ inhibitors due to variations in the platelet inhibitory effect and pharmacodynamic and pharmacokinetic properties.⁶⁷ In the CABG substudy of the Clopidogrel in Unstable Angina to Prevent Recurrent Events (CURE) trial, > 5 days' discontinuation of clopidogrel before surgery did not increase the risk of bleeding [risk ratio (RR) 0.83, 95% CI 0.46–1.48].⁶¹ For prasugrel, a time interval of 7 days is recommended due to the longer offset time compared with clopidogrel⁶⁸ and the high incidence of bleeding complications reported in the CABG substudy of the Trial to Assess Improvement in Therapeutic Outcomes by Optimizing Platelet Inhibition with Prasugrel–Thrombolysis in Myocardial Infarction (TRITON-TIMI) 38 trial.³⁶ Although it was previously recommended that ticagrelor be discontinued at least 5 days before surgery,⁶⁹ a large observational study in CABG patients showed that discontinuation of ticagrelor for 3 or 4 days before surgery did not relate to a higher incidence of bleeding complications (OR 0.93, 95% CI 0.53–1.64)³⁷; this observation is also supported by others.^{38,63} Current guidelines recommend restarting DAPT in all patients with ACS, independent of the revascularization strategy.^{66,67}

Because of the individual variation in the magnitude and duration of the antiplatelet effect,^{21,69,70} the use of platelet function tests may help to optimize the timing of surgery or

establish the grade of platelet inhibition in patients in whom the time since discontinuation is unclear. The guidance is that the interruption of therapy may be preferred over an arbitrary, specified period.^{66,67} The platelet inhibitory response assessed by platelet function testing to clopidogrel,^{22,32,33,71} prasugrel^{22,32,33,71} and ticagrelor²¹ is associated with CABG-related bleeding. A strategy based on preoperative platelet function testing to determine the timing of CABG in clopidogrel-treated patients led to 50% shorter waiting times for surgical treatment compared to those for a discontinuation time-based strategy.²⁰ It should, however, be pointed out that no RCT or observational study has investigated the incidence of perioperative bleeding complications in relation to the time since discontinuation-based or platelet function test-based surgery and that defined cut-off levels for acceptable platelet function are unavailable.

Taken together, in patients taking DAPT who need to undergo non-emergent cardiac surgery, postponement of surgery for at least 3 days after discontinuation of ticagrelor, 5 days after clopidogrel and 7 days after prasugrel should be considered to reduce the risk of postoperative bleeding (Fig. 1). During discontinuation of P2Y₁₂ inhibitors, ASA therapy should be continued until surgery. Platelet function testing may be used to guide the timing of surgery in patients taking DAPT.

4.2.3. Glycoprotein IIb/IIIa Inhibitors

Background. Glycoprotein IIb/IIIa (GPIIb/IIIa) inhibitors (eptifibatide, tirofiban and abciximab) are today almost exclusively used in conjunction with percutaneous coronary interventions but may also be used for bridging risk patients on oral P2Y₁₂ inhibitors to surgery.^{66,67,72}

Description of the Evidence. The optimal time for discontinuation of GPIIb/IIIa inhibitors before surgery is mainly based on pharmacokinetic calculations. Platelet function recovery is obtained 24–48 h after discontinuation of abciximab and 4–8 h after discontinuation of eptifibatide and tirofiban.⁷³ Cessation at 4 h before surgery is sufficient for patients treated with GPIIb/IIIa inhibitors. In a small prospective study, discontinuation of eptifibatide 4 h before CABG resulted in lower bleeding rates than discontinuation 2 h before the procedure.⁷⁴ For abciximab, no difference in bleeding was noted in a pooled analysis of 82 patients in the Evaluation in Percutaneous Transluminal Coronary Angioplasty to Improve Long-Term Outcome with Abciximab GP IIb/IIIa Blockade (EPILOG) and Evaluation of Platelet IIb/IIIa Inhibitor for Stenting (EPISTENT) trials who underwent acute CABG when abciximab was stopped < 6 h before CABG in 61% of the patients.⁷⁵ In a small retrospective study, tirofiban-treated patients having CABG showed more bleeding than patients without tirofiban, but no difference was found between different discontinuation times.⁷⁶

In conclusion, GPIIb/IIIa inhibitor discontinuation at least 4 h before surgery should be considered to minimize the risk of post-operative bleeding.

4.2.4. Low-Molecular-Weight Heparin

Background. Low-molecular-weight heparin (LMWH; enoxaparin and fondaparinux) mainly inhibits activated FX (FXa), with plasma concentration peaks 3–4 h after administration. It is the preferred strategy for prophylactic and therapeutic anticoagulation in patients after certain surgeries (e.g. orthopaedic surgery) and in patients with malignancies. The half-life is around 5 h in patients with normal renal function. Monitoring of the anticoagulant effects of LMWH can be achieved by plasma anti-FXa activity. LMWH-induced bleeding may be treated with protamine, but this therapy does not completely reverse the anticoagulant effect of LMWH.

Description of the Evidence. Cardiac surgery patients receiving enoxaparin versus unfractionated heparin (UFH) more frequently underwent re-exploration for bleeding (HR 2.6, 95% CI 1.1–5.9).⁴⁴ Patients receiving LMWH within 12 h before surgery showed more blood loss and transfusion requirements than patients receiving UFH or a dose of LMWH > 12 h before surgery.⁷⁷ However, other studies could not confirm this finding.⁴¹ Fondaparinux versus enoxaparin before CABG surgery resulted in the same rate of bleeding events, but when it was not discontinued 36 h before surgery, fondaparinux was associated with higher 12-h postoperative bleeding rates.⁴² The half-life of fondaparinux is increased in patients with chronic kidney disease.

Overall, preoperative bridging of oral anticoagulation with UFH/LMWH is only indicated in patients at high risk of thrombotic events. It is further recommended that prophylactic LMWH be discontinued 12 h and fondaparinux 24 h before surgery; a longer interval may be necessary for patients with impaired renal function.

4.2.5. Vitamin K Antagonists

Background. Vitamin K antagonists (VKAs) are commonly used to prevent and treat thromboembolism in patients with atrial fibrillation, venous thromboembolic disease and mechanical heart valves (particularly when they are in the mitral or tricuspid position). The anti-coagulation effect can be monitored by the prothrombin time and is most often expressed as an international normalized ratio (INR).

Description of the Evidence. VKAs are regularly stopped 3–5 days before surgery to obtain an INR < 1.5. In patients having urgent or emergency surgery, the effect of VKA can be completely reversed by administering prothrombin complex concentrate (PCC). Bridging non-cardiac surgery patients who are taking VKA with a full therapeutic dose of LMWH after surgery is associated with increased bleeding but not with a meaningful reduction in thrombotic events.³⁹ Thus, bridging to cardiac surgery is only recommended in patients at a very high risk of thrombotic events, e.g. those with a recent (< 4 weeks) pulmonary embolism or a prosthetic mechanical valve, atrial fibrillation with the rheumatic valvular disease or atrial fibrillation with a CHA₂DS₂-VASc [congestive heart failure,

hypertension, age ≥ 75 (2 points), diabetes, prior stroke (2 points) – vascular disease, age 65–74, sex category (female)] score > 4. In these patients, VKA should be discontinued 5 days before surgery and bridged by LMWH (Fig. 2). Bridging with UFH after surgery until the INR reaches the therapeutic range in patients on VKAs (bridging with subcutaneous LMWH may be considered an alternative to bridging with UFH) is recommended. Elective cardiac surgery should not be performed if the INR is > 1.5 in patients taking VKAs. When surgery cannot be postponed, coagulation factors should be used to reverse the effect.

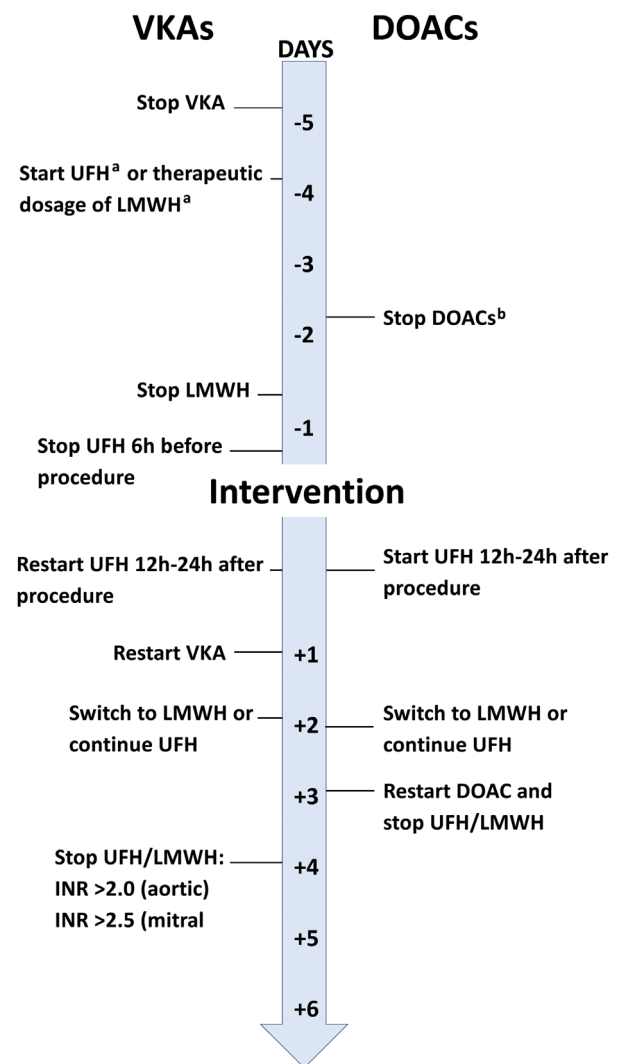


Figure 2. Management of oral anticoagulation in patients with an indication for pre- and/or postoperative bridging (reproduced with permission from Sousa-Uva, Stuart J. Head, Milan Milojevic, Jean-Philippe Collet, Giovanni Landoni, Manuel Castella et al. 2017 EACTS Guidelines on Perioperative Medication in Adult Cardiac Surgery. *Eur J Cardiothorac Surg* 2017; <http://dx.doi.org/10.1093/ejcts/ezx314>). ^aBridging with UFH/LMWH should start when INR values are below specific therapeutic ranges. ^bDiscontinuation should be prolonged to > 72 h if creatinine clearance is 50–79 ml/min/1.73 m² or ≥ 96 h if creatinine clearance is < 50 ml/min/1.73 m². DOACs: direct oral anticoagulants; INR: international normalized ratio; LMWH: low-molecular-weight heparin; UFH: unfractionated heparin; VKAs: vitamin K antagonists.

4.2.6. Direct Oral Anticoagulant

Background. Direct oral anticoagulants (DOACs) include direct thrombin inhibitors (dabigatran) and FXa inhibitors (rivaroxaban, apixaban and edoxaban), and novel formulations are under development. DOACs are increasingly used as an alternative anticoagulation strategy for VKAs.

Description of the Evidence. Because emergency surgery in patients under dabigatran treatment has been associated with severe or even fatal bleeding, it is recommended that DOACs be discontinued at least 48 h before cardiac surgery.^{78–80} DOACs may be bridged by heparins in patients with acute thrombotic events < 4 weeks. The half-life of DOACs may be prolonged in case of impaired renal function (Table 3).

Whenever feasible in patients with impaired renal function, the concentration of DOAC should be measured in terms of the diluted thrombin time (dabigatran) or calibrated with anti-FXa activities (rivaroxaban, apixaban or edoxaban). The use of viscoelastic tests is still debated in the diagnosis of DOAC-induced prolonged clotting time. Ecarin clotting times may be used when available for patients taking dabigatran. For emergency reversal of dabigatran, the newly released antidote (idarucizumab) can be used in the pre- and postoperative settings. Treatment of post-operative FXa-related bleeding includes PCC, activated PCC (FEI-BAVR, Shire US Inc., Lexington, MA, USA) and recombinant activated factor VII (rFVIIa), because no specific antidote is approved at the moment. These agents have been found to be effective in some studies (PCC in volunteer studies, aPCC and rFVIIa mainly *in vitro* studies) on FXa inhibitors. Due to the relatively low level of protein binding, haemodialysis is effective for increasing the clearance of dabigatran but not for that of FXa-inhibitors (protein binding > 85%).

Based on the available evidence, it is recommended that DOACs be stopped at least 48 h prior to surgery in patients having elective cardiac surgery; a longer interval may be necessary for patients with impaired renal function (Fig. 2).

Table 3
Different types of DOACs

	Rivaroxaban	Apixaban	Edoxaban	Dabigatran
Target	Factor Xa	Factor Xa	Factor Xa	Thrombin
T _{max}	2–4 hours	3 hours	1–3 hours	2 hours
Half-life	9–13 hours	9–14 hours	5–11 hours	14–17 hours
Frequency	Once daily	Twice daily	Once daily	Once or twice daily
Renal excretion	66% (half inactive)	25%	36–45%	80%
Antidote	Andexanet ^a	Andexanet ^a	Andexanet ^a	Idarucizumab
Discontinuation	48 hours ^b	48 hours	48 hours	48–96 hours ^b

EMA: European Medicines Agency; DOACs: direct oral anticoagulants.

^aNot yet approved by EMA.

^bDiscontinue > 48 h if creatinine clearance is > 80 ml/min/1.73 m²; discontinue > 72 h if creatinine clearance is 50–79 ml/min/1.73 m²; and discontinue > 96 h if creatinine clearance is < 50 ml/min/1.73 m².

4.3. Preoperative Anaemia

Recommendations	Class ^a	Level ^b	Ref ^c
Oral or intravenous iron alone prior to cardiac surgery may be considered in mildly anaemic patients (women, Hb 100–120 g/l; men, Hb 100–130 g/l) or in severely anaemic patients (both genders, Hb ≤ 100 g/l) to improve erythropoiesis.	IIb	C	
Erythropoietin with iron supplementation should be considered to reduce postoperative transfusions in patients with non-iron deficiency (e.g. EPO, vitamin D or folate acid deficiency) undergoing elective surgery.	IIa	B	81–83
Preoperative erythrocyte transfusion is not routinely recommended in preoperative anaemic patients to prevent postoperative AKI.	III	C	

^aClass of recommendation.

^bLevel of evidence.

^cReferences.

AKI: acute kidney injury; EPO: erythropoietin; Hb: haemoglobin.

4.3.1. Implications for Preoperative Anaemia

Background. Anaemia is a frequent comorbid condition in the elderly patients, and up to 40% of patients present with mild anaemia (women, Hb 100–120 g/l; men, Hb 100–130 g/l) or severe anaemia (both genders, Hb ≤ 100 g/l) of any cause prior to cardiac surgery.⁸⁴ Although anaemia is recognized to be a predictor of postoperative adverse outcomes, this evidence is mostly based on observational studies.

Description of the Evidence. Preoperative anaemia is associated with a worsening of the clinical outcome after cardiac surgery in terms of blood transfusion requirements, acute kidney injury (AKI) and death.^{85–87} Several retrospective studies showed a significant interaction between preoperative anaemia and blood transfusions on outcome and a pronounced detrimental effect of blood transfusions for anaemic patients.^{88,89} The risk for intraoperative transfusion is significantly higher in mildly anaemic (HR 1.44, 95% CI 1.20–1.73) or severely anaemic patients (HR 1.81, 95% CI 1.34–2.44).^{88,89}

Derived from retrospective studies, preoperative anaemia is, therefore, considered a risk factor for adverse clinical outcomes and transfusion requirements after cardiac surgery.

4.3.2. Iron Supplementation

Background. Supplementation of oral or intravenous iron is the treatment of choice for iron deficiency anaemia. Preoperative iron supplementation may reduce transfusion requirements

in patients undergoing cardiac surgery with iron deficiency anaemia.

Description of the Evidence. A recent RCT in non-anaemic patients, based on a 3-arm parallel group design, compared the effect of 3 preoperative doses of intravenous iron sucrose (100 mg) and daily oral iron (125 mg) with placebo until 1 month following discharge. No difference was found between the transfusion requirements of the various groups.⁹⁰ The PROphylaxis for ThromboEmbolism in Critical Care Trial (PROTECT) investigated the perioperative administration of iron isomaltoside (1000 mg) on postoperative Hb levels and transfusion requirements in non-anaemic patients having elective surgery.⁹¹ Although there was a lower number of non-anaemic patients in the iron isomaltoside-treated group compared with the group receiving a placebo 1 month after surgery (8.0% vs 38.5%, $P = 0.019$), transfusion requirements up to 4 weeks after surgery were not different between groups (13.3% vs 20.0%, $P = 0.52$). Apart from the significant effect of iron supplementation on stimulation of erythropoiesis, there are limited data supporting the association between the administration of iron and reduced postoperative transfusion requirements in patients with anaemia undergoing cardiac surgery. The optimal time to start iron therapy before surgery is unclear from the current evidence.

4.3.3. Erythropoietin

Background. Preoperative anaemia is associated with adverse outcomes after cardiac surgery in several retrospective studies.^{92,93} Erythropoietin (EPO) with or without iron supplementation may be used to treat preoperative anaemia and to reduce the risk of postoperative transfusion requirements.

Description of the Evidence. Erythropoietin, with or without iron, has been used to treat noniron deficiency anaemia and to increase red cell mass in patients who refuse blood transfusions or who are at high risk of postoperative anaemia, such as patients with renal failure.^{94,95} Administration of EPO (80 000 IU) and oral iron (40 mg) or iron alone in 600 mildly or non-anaemic patients reduced the perioperative PRBC transfusion requirements in patients treated with EPO more than in controls (17% vs 39%, $P < 0.0005$), but this difference was diminished after controlling for baseline Hb levels.⁸¹ In 320 patients undergoing off-pump CABG surgery, preoperative high-dose EPO treatment (cumulative dose of 52 000 IU) starting 2 days prior to and lasting until 2 days after surgery was compared to the effects of placebo.⁸² Postoperative transfusion requirements were lower in the EPO group compared with the placebo group (0.33 vs 0.76 units per patient, $P = 0.008$). A single dose of EPO (500 IU/kg) with iron sucrose (200 mg) 1 day prior to surgery in primary heart valve procedures reduced transfusion requirements during surgery until 4 days postoperatively compared with the control group (59.5% vs 86.5%, $P = 0.009$).⁸³ In all studies, the investigators were blinded to the study treatment (single blinded).

In summary, EPO with or without iron supplementation should be considered in patients with non-iron deficiency undergoing elective surgery to reduce postoperative transfusions. Even an EPO infusion in the days before surgery may have benefits in terms of transfusion requirements. It should be noted that, in part, data showing a treatment benefit for EPO were obtained from non-anaemic treated patients undergoing cardiac surgery.

4.3.4. Blood Transfusion to Treat Preoperative Anaemia

Background. Preoperative blood transfusions to treat anaemia are frequently used to symptomatically treat low Hb values. Transfusion prior to cardiac surgery or with initiation of CPB has been suggested for patients with pre-existing anaemia.

Description of the Evidence. One small RCT investigated the effect of preoperative PRBC transfusion in anaemic patients on intraoperative transfusion requirements and AKI as outcomes.⁹⁶ Prophylactic transfusion of 2 PRBC units was associated with a reduced rate of intraoperative transfusion [0 (0–2) vs 2 (1–4) units, $P < 0.001$] compared with standard care but not with a reduced rate of AKI.⁹⁶ Given the lack of additional published evidence, an expert consensus was reached by the guideline committee against the routine use of transfusions for patients with preoperative anaemia. However, in the case of emergency surgery and life-threatening anaemia, it is legitimate to use preoperative blood transfusions to increase Hb levels.

5. Intraoperative Management

Intraoperative preservation of patient haemostasis is a multidisciplinary and multifactorial challenge. Even though the surgeon applies meticulous haemostasis and patience with respect to clot formation, these measures are only effective when paralleled by interventions that minimize haemodilution, normothermia, appropriate anticoagulation and haemostatic monitoring during the procedure.

5.1. Surgical Techniques

Special surgical or perfusion techniques, such as off-pump CABG, minimally invasive extracorporeal circulation circuits (MiECC) and minimally invasive cardiac surgery are not universally implemented. These procedures might only be safe and efficiently performed when integrated into the daily clinical routine. Therefore, any recommendations regarding the use of these techniques are addressed to centres with adequate experience in these procedures. However, especially for severely anaemic patients, it is highly recommended that the members of the multidisciplinary team (cardiologists, surgeons, anaesthesiologists and perfusionists) discuss optimal treatment strategies, including surgical techniques, the limitations of haemodilution and improved CPB systems to avoid massive transfusions and bleeding.

Recommendations	Class ^a	Level ^b	Ref ^c
It is recommended that the members of the multidisciplinary team ^d discuss the optimal surgical strategy based on clinical status, comorbidities, bleeding risk and team expertise.	I	C	
Off-pump CABG surgery may be considered in selected patients ^e to reduce perioperative transfusions.	Iib	B	97,98
MiECC systems should be considered over standard conventional CPB systems to reduce perioperative transfusions.	Iia	B	99,100
Minimally invasive heart valve surgery may be considered to reduce blood loss and the need for transfusions.	Iib	B	101–103
Routine use of topical sealants in cardiac surgery is not recommended.	III	B	104
Topical sealants may be considered in clinical situations where conventional approaches to surgical and medical improvement of haemostasis and where bleeding problems are more local than generalized.	Iib	C	

^aClass of recommendation.

^bLevel of evidence.

^cReferences.

^dSurgeons, cardiologists, anaesthesiologists and perfusionists.

^eDialysis-dependent patients, patients operated on under DAPT and anaemic patients with low body surface area.

CABG: coronary artery bypass grafting; CPB: cardiopulmonary bypass; DAPT: dual antiplatelet therapy; MiECC: minimally invasive extracorporeal circulation circuits.

5.1.1. Off-pump Surgery

Background. Several hypothetical mechanisms by which off-pump CABG surgery might reduce bleeding and/or the need for blood product transfusions, including lower dosing of systemic heparin, the standard use of cell saving and the fact that haemodilution and blood trauma caused by CPB, are obviated with this technique.

Description of the Evidence. There are no RCTs comparing off-pump and on-pump CABG where bleeding, transfusion or the need for reoperation was the primary end point; the data are therefore extrapolated. A meta-analysis comprising 102 RCTs compared off-pump with on-pump CABG in 19 101 patients.⁹⁸ The secondary end points of PRBC transfusion (OR 0.49, 95% CI 0.33–0.72) and blood product transfusion (OR 0.66, 95% CI 0.55–0.81) were significantly lower in patients having off-pump than in those having on-pump CABG surgery, without differences in re-exploration for

bleeding.⁹⁸ However, a large variability across studies was observed, suggesting that such a benefit could not be reproduced in several of the RCTs included. A recent meta-analysis of RCTs showed similar results⁹⁷ but was also confounded by a high grade of heterogeneity for transfusion requirements and chest tube drainage among RCTs. In the 2 largest RCTs (the Coronary Artery Bypass Surgery [CABG] Off or On Pump Revascularization Study [CORONARY]¹⁰⁵ and the German Off-Pump Coronary Artery Bypass Grafting in Elderly Patients [GOPCABE] trial¹⁰⁶), off-pump surgery was associated with lower transfusion rates (50.7% vs 63.3% and 56.3% vs 62.7%, respectively; both $P < 0.001$). However, in these RCTs, physicians were not blinded to treatment allocation, and the transfusion protocol was not prespecified. Therefore, the treatment received might have influenced the decision to transfuse.

In summary, off-pump surgery may be considered a surgical technique that is associated with fewer transfusion requirements than on-pump surgery.

5.1.2. Minimally Invasive Extracorporeal Circulation

Background. The common features of MiECC are a small priming volume (~600–750 ml), a reduced artificial surface due to the absence of the venous reservoir, a biocompatible coated system and a centrifugal pump. Systems are completely closed to avoid blood–air contact, and they include the separation of shed blood.¹⁰⁷

Description of the Evidence. RCTs comparing MiECC versus conventional CPB were too underpowered to detect a significant difference in transfusion rates. The available meta-analyses^{99,100} show a reduced rate of transfusion with MiECC but include open studies and studies without a predefined transfusion protocol. The type of circuit used may also have influenced the decision to transfuse. These aspects limit the generalizability of current comparisons. Two recent meta-analyses compared MiECC with conventional CPB procedures.^{99,100} An analysis of the results of 24 studies ($n = 2770$) showed that the use of MiECC was associated with a reduced risk of PRBC transfusion (OR 0.24, 95% CI 0.16–0.37; $P < 0.001$) but without a reduction in reoperations for bleeding.⁹⁹ In a meta-analysis that assessed 29 studies with 2335 patients, the use of MiECC resulted in a significant reduction in the risk of transfusions (OR 0.35, 95% CI 0.23–0.53), whereas reoperation rates for persistent bleeding revealed no difference.¹⁰⁰

In summary, the use of MiECC systems, when compared with conventional CPB systems, is associated with a reduction in PRBC transfusions. Therefore, in an attempt to reduce allogeneic blood transfusions, MiECC systems can be considered. However, the use of MiECC systems does not contribute towards reducing reoperation rates for bleeding.

5.1.3. Minimally Invasive Surgery

Background. Minimally invasive heart valve surgery may play a role in reducing the need for transfusion and reoperation

for bleeding, because it requires a smaller incision and less tissue dissection and retraction to open up the surgical field.

Description of the Evidence. The available evidence consists almost entirely of observational studies and should not be considered definitive. The 2010 International Society for Minimally Invasive Cardiothoracic Surgery consensus statement concluded that the risk of transfusion for PRBCs was reduced after minimal mitral valve surgery compared with the conventional approach (difference of -1.9 units; 95% CI -2.5 to -1.2 units), but reoperation for bleeding was not different between the groups.¹⁰² A systematic review of mitral valve surgery through a right lateral minithoracotomy versus sternotomy identified a total of 45 studies, including 3 RCTs.¹⁰¹ The minimally invasive approach was associated with a small reduction in bleeding (mean difference -142.1 ml, 95% CI -199.2 to -85.1 ml) and risk for transfusion (RR 0.67, 95% CI 0.51–0.88). However, pooled data from the small RCTs showed that minimally invasive surgery did not reduce total blood loss (mean difference -149 ml; 95% CI -491 to 193 ml) or the incidence of transfusion (RR 0.94, 95% CI 0.61–1.45). In a systematic review on aortic valve replacement through minimally invasive access versus full sternotomy,¹⁰³ pooled data from 3 RCTs showed no difference in the rate of blood product transfusions (RR 0.77, 95% CI 0.58–1.03), whereas pooled data from observational studies suggested otherwise. Minimally invasive heart valve surgery is practised in high volumes in only a limited number of centres.

Until minimally invasive valve surgery becomes more widely adopted and large RCTs are performed, there is not enough evidence to suggest broad adaption for the purpose of blood conservation.

5.1.4. Topical Haemostatics

Background. In addition to systemic haemostatic action, topical haemostatic agents are locally applied and include haemostatic sealants and topical antifibrinolytic agents [aprotinin, tranexamic acid (TXA) and E-aminocaproic acid (EACA)].^{108,109} Passive haemostatic agents that activate platelets, including collagen, gelatines and regenerated oxidized cellulose, do not contain biologically active clotting factors. The last category of topical agents includes synthetic (cyanoacrylates) and semi-synthetic (glutaraldehyde-albumin) sealants, which have sealing properties through a rapid polymerization process without having intrinsic haemostatic activity.

Description of the Evidence. A Cochrane systematic review¹⁰⁸ on 18 RCTs comparing fibrin sealant versus no fibrin sealant concluded that fibrin sealants are efficacious in reducing both postoperative blood loss (mean difference -161 ml, 95% CI -98.25 to -224.53) and perioperative exposure to allogeneic PRBC transfusions (RR 0.63, 95% CI 0.45–0.88). However, this review was limited by the inclusion of only 1 cardiac surgery study and RCTs with small sample sizes. A meta-analysis¹⁰⁹ including 8 RCTs ($n = 622$ patients) showed that topical antifibrinolytic agents reduced the amount of 24-h chest tube blood loss by 220 ml (95% CI -318 to -126) and

resulted in a saving of 1 PRBC unit per patient (95% CI -1.54 to -0.53). However, high heterogeneity among RCTs was found. No adverse effects were reported following topical use of the medications. A more recent Cochrane systematic review¹¹⁰ including 7 RCTs ($n = 511$) supported a protective role of topical antifibrinolytic agents in cardiac surgery (RR 0.63, 95% CI 0.61–0.66), but the overall quality of the studies included was low. A recent RCT including off-pump CABG patients did not find any benefit from topical EACA regarding postoperative bleeding and transfusion.¹¹¹

Based on the available evidence, the routine use of topical sealants in cardiac surgery is not recommended and may only be considered in cases of persistent bleeding where the bleeding is localized.

5.2. Cardiopulmonary Bypass

Recommendations	Class ^a	Level ^b	Ref ^c
Implementation of institutional measures to reduce haemodilution by fluid infusion and CPB during cardiac surgery to reduce the risk of bleeding and the need for transfusions is recommended.	I	C	
The use of a closed extracorporeal circuit may be considered to reduce bleeding and transfusions.	IIb	B	112,113
The use of a biocompatible coating to reduce perioperative bleeding and transfusions may be considered.	IIb	B	114–116
The routine use of cell salvage should be considered to prevent transfusions.	IIa	B	117–119
(Modified) ultrafiltration may be considered as part of a blood conservation strategy to minimize haemodilution.	IIb	B	120–122
Retrograde and antegrade autologous priming should be considered as part of a blood conservation strategy to reduce transfusions.	IIa	A	123–125
Normothermia during CPB (temperature > 36° C) and maintenance of a normal pH (7.35–7.45) may contribute to a reduced risk of postoperative bleeding.	IIb	B	126,127

^aClass of recommendation.

^bLevel of evidence.

^cReferences.

CPB: cardiopulmonary bypass.

5.2.1. Closed versus Open Cardiopulmonary Bypass

Background. The contact of blood with ambient air in the reservoir and in other parts of the circuit in extracorporeal circulation (ECC) is regarded as one of the several causes of

coagulation factor activation during CPB.¹²⁸ Closed systems minimize the blood/air interface and platelet activation and are an integral part of MiECC systems but result in more complicated perfusion that requires active trapping of air bubbles.¹²⁹ The use of cardiotomy suction requires a separate reservoir, and vacuum-assisted venous drainage cannot be applied to the collapsible reservoir bag.

Description of the Evidence. A large number of studies combined several components of blood conservation, including closed circuits. Some studies combined the elimination of cardiotomy suction with the use of a closed system or investigated the combined effect of surface coating, a closed system and different pumps.^{113,130} In an RCT, closed systems had no advantage over open systems for patient haemostasis.¹¹³ In a small, single-centre RCT, a closed system reduced transfusion requirements compared with an open system (378 ± 364 ml vs 717 ± 486 ml; $P = 0.003$).¹¹² Taken together, the use of a closed ECC may be considered to reduce bleeding and transfusions, but adequately sized RCTs on closed versus open systems are lacking.

5.2.2. Biocompatible Coating

Background. Biocompatible coating of tubing and oxygenator mimics the natural endothelial lining of blood vessels and improves the haemocompatibility and hydrophilicity of the system, thereby reducing the risk of clot activation. Among others, biocompatible coatings include ionic or covalent heparin bonding, poly(2-methoxyethylacrylate) (PMEA) and phosphorylcholine. The effect of the biocompatible coating of the ECC on bleeding and transfusion requirements is still under investigation.

Description of the Evidence. A systematic review and meta-analysis published in 2009 of 36 RCTs issued between 1992 and 2006 showed that the use of any biocompatible coating reduced the risk of PRBC transfusion (OR 0.8, 95% CI 0.69–0.93) compared with the use of a non-coated circuit.¹¹⁴ A more recent systematic review of 14 RCTs concluded that second and third-generation heparin-coated circuits only showed superiority on perioperative blood loss to non-coated circuits in about 50% of the studies included.¹¹⁶ A large RCT showed that the combination of heparin-coated circuits with low or conventional dose heparinization was associated with a reduced 12-h blood loss compared with non-coated circuits (mean volume 382 vs 431 vs 457 ml; $P < 0.01$, respectively).¹³¹

Other studies showed that the use of phosphorylcholine-coated circuits might contribute to less perioperative blood loss and fewer transfusion requirements.^{132–134} In 2 small studies comparing the effect of PMEA, phosphorylcholine and heparin-coated circuits,^{135,136} only 1 study showed that PMEA coating was associated with fewer platelet transfusions compared with heparin coating.¹³⁵

Based on the available data, the use of a biocompatible coating may be considered as part of a PBM programme to reduce perioperative bleeding and transfusions.

5.2.3. Cell Salvage

Background. Cell salvage of operative blood loss and residual blood from the circuit after CPB may contribute to a reduction in the use of allogeneic blood products, especially PRBCs. A specific rationale for cell salvage is its use during CPB instead of cardiotomy suction, since the retransfusion of whole blood using cardiotomy suction through the ECC may contribute to systemic inflammation and bleeding.

Description of the Evidence. A Cochrane review showed that the use of cell salvage reduced the risk of PRBC transfusions (RR 0.78, 95% CI 0.68–0.91) during cardiac surgical procedures.^{117,137} In a meta-analysis comparing cell salvage with cardiotomy suction, it was found that cell salvage reduced the risk of any blood transfusion (OR 0.63, 95% CI 0.43–0.94) without increasing adverse effects or bleeding.¹³⁸ Recent studies confirmed that the use of cell salvage is efficient in reducing allogeneic blood exposure.^{118,119,139}

Cardiotomy suction might be associated with thrombin generation and neutrophil and platelet activation and could contribute to higher transfusion requirements.⁹ The amelioration of inflammation by the removal of activated plasma through the washing of shed blood has repeatedly been demonstrated.^{140–142} Cell saver processing of cardiotomy suction blood only has no beneficial effect on blood conservation and increases the use of fresh-frozen plasma (FFP) transfusions.^{138,143,144} Extending the use of cell salvage to the postoperative phase may show additional benefits,¹⁴³ although others did not confirm this.¹¹⁸ Moreover, large volumes of cell-salvaged blood may lead to a relative loss of plasma and may result in impaired coagulation.¹⁴⁵ Therefore, it may be concluded that the balance between a positive effect of cell salvage on transfusion requirements and a reduction of negative systemic effects (versus the inherent loss of plasma, platelets and leucocytes) depends on the quantity of the salvaged blood.

In the postoperative period, tube drainage and the washing of blood may be implemented to reduce postoperative blood transfusions. The British National Institute for Health and Clinical Excellence conducted an extensive review of the literature and several meta-analyses for postoperative cell salvage.¹⁴⁶ In cardiac surgery, the absolute risk of being exposed to allogeneic blood was approximately 15.6% (95% CI 7.4–21.4%) lower in patients receiving postoperative cell salvage and 1 unit less of allogeneic blood. Postoperative cell salvage resulted in lower rates of infection, lower mortality rates and shorter hospital stays. Unwashed salvaged blood from postoperative surgical drains contains higher concentrations of inflammatory mediators, fibrin split products, interleukins, fat emboli and complement factors and may increase the risk of inflammatory complications.¹⁴⁷

In summary, the routine use of cell salvage should be considered to prevent transfusions, but the retransfusion of large volumes of cell salvaged blood (> 1000 ml) may impair coagulation. Overall, postoperative cell salvage and reinfusion of washed erythrocytes may be considered to reduce transfusions in patients with bleeding.

5.2.4. Ultrafiltration

Background. The total volume of blood during CPB can be controlled using ultrafiltration. In modified ultrafiltration, excess fluid is discarded from the remaining blood after the termination of ECC.

Description of the Evidence. Ultrafiltration and modified ultrafiltration are accompanied by removal of mediators of inflammation, which may also lead to improved haemostasis. In a meta-analysis, it was shown that ultrafiltration, and particularly modified ultrafiltration, was associated with a reduction in postoperative blood transfusions (-0.73 units; 95% CI -1.16 to -0.31).¹²¹ In a large RCT, patients subjected to ultrafiltration showed lower transfusion requirements compared to patients in whom ultrafiltration was not implemented.¹²²

Although most studies hinted at a beneficial effect of ultrafiltration on postoperative transfusion requirements, especially in the presence of preoperative anaemia, the evidence is based on small underpowered studies, with transfusion requirements as a secondary end point.

5.2.5. Retrograde and Antegrade Autologous Priming

Background. The priming of the ECC with an asanguineous fluid results in haemodilution and increases the risk of blood transfusion, particularly in patients with a small body surface area.¹⁴⁸ Antegrade autologous priming and retrograde autologous priming (RAP) are simple, inexpensive, efficient ways to address the issue of haemodilution.

Description of the Evidence. Most of the available literature on this topic refers to studies in which several features of minimized bypass systems were combined. However, some smaller RCTs focusing on RAP have been performed. The largest meta-analysis concluded that RAP significantly reduced transfusion requirements (RR 0.53, 95% CI 0.43–0.66; $P < 0.001$).¹²³ Similar findings were demonstrated in the meta-analysis of 6 RCTs in which patients treated with RAP received significantly fewer PRBC transfusions during hospitalization (weighted mean difference -0.60 units, 95% CI -0.90 to -0.31; $P = 0.001$).¹²⁴ These findings were confirmed in an RCT with 120 patients and a body surface area $< 1.5 \text{ m}^2$, showing that RAP resulted in a marked reduction of haemodilution and significantly reduced transfusion requirements.¹⁴⁸ On the basis of the evidence, RAP and antegrade autologous priming should be considered as part of a blood conservation strategy to reduce transfusions.

5.2.6. Coagulation-Friendly Environment

Background. The enzymatic process of coagulation requires an optimal temperature and acidity level. Both hypothermia and acidosis may reduce the thrombin-generation capacity, thereby impairing patient haemostasis. The number of studies focusing on the association of hypothermia or acidosis with postoperative bleeding is limited.

Description of the Evidence. Hypothermia in cardiac surgery is used to enhance myocardial and end-organ preservation, particularly during CPB. However, hypothermia is also

associated with coagulopathy. In an RCT of patients who had CABG surgery, it was shown that hypothermia (34°C) was not associated with greater 12-h blood loss compared with normothermic patients (37°C).¹⁴⁹ A retrospective analysis of patients having off-pump CABG surgery showed that transfusion rates were higher in patients exposed to mild hypothermia ($< 36^\circ \text{C}$).¹²⁷ One retrospective study investigated the association between acidosis, post-surgery acidosis and postoperative bleeding,¹²⁶ showing that even a moderate degree of acidosis ($\text{pH} < 7.35$) and hyperlactataemia (lactate $> 4.0 \text{ mmol/l}$) was associated with a significantly higher volume of postoperative chest drainage compared with no acidosis/hyperlactataemia 12 h after surgery (mean blood loss 576 vs 406 ml; $P = 0.001$).

Despite the lack of large, prospective studies on temperature management and the maintenance of pH during cardiac surgery, there is general agreement that the coagulation system requires normal physiological circumstances, including normothermia and a pH nearly 7.4.

5.3. Intraoperative Anticoagulation

Recommendations	Class ^a	Level ^b	Ref ^c
Heparin level-guided heparin management should be considered over ACT-guided heparin management to reduce bleeding.	IIa	B	150–152
Heparin level-guided protamine dosing may be considered to reduce bleeding and transfusions.	IIb	B	153
Protamine should be administered in a protamine-to-heparin dosing ratio ^d $< 1:1$ to reduce bleeding.	IIa	B	154
AT supplementation is indicated in patients with AT deficiency to improve heparin sensitivity.	I	B	155,156
AT supplementation is not recommended to reduce bleeding following CPB.	III	C	
In patients with HIT antibodies for whom surgery cannot be postponed, anticoagulation with bivalirudin should be considered when the bleeding risk is acceptable. The use of heparin in the pre- and postoperative periods should be avoided.	IIa	C	

^aClass of recommendation.

^bLevel of evidence.

^cReferences.

^dProtamine-to-heparin dosing ratio based on the initial heparin dose.

ACT: activated clotting time; AT: antithrombin; CPB: cardiopulmonary bypass; HIT: heparin-induced thrombocytopenia.

5.3.1. Heparin and Anticoagulation Monitoring

Background. UFH binds to antithrombin (AT), which then potentiates the inactivation of thrombin and FXa by AT up to a

1000-fold. The sensitivity to heparin is determined by patient-specific characteristics¹⁵⁷ and is assessed by the whole-blood activated clotting time (ACT) test. Target ACT values range from 300 to 600 s depending on the methods used to measure the ACT or due to different heparin dosing strategies. Moreover, the variation in the efficacy of different heparins may need individual anticoagulation strategies.

UFH is usually dosed based on patient body weight (300–600 U/kg), followed by additional doses in the case of a periprocedural decrease in the ACT. Heparin resistance and postoperative heparin rebound are major limitations of this strategy. Heparin resistance may lead to insufficient anticoagulation during the procedure, whereas heparin rebound may contribute to post-surgical bleeding.

Description of the Evidence

Individualized Heparin Management. Individualized heparin management or titration based on a dose–response test is based on the use of the HMS/HepCon (Medtronic, Minneapolis, MN, USA), Hemochron RxDx (Accriva Diagnostics, San Diego, CA, USA) or anti-Xa measurements in addition to the ACT. The number of RCTs that focus on the advantages of heparin dose–response implementation is, however, limited. In most studies, the use of a heparin titration device resulted in an increased heparin dose and a decrease in the protamine dose. In most studies, this situation is associated with decreased blood loss and transfusion requirements as well as with higher platelet counts at the end of the operation.^{150,151,153,158–160} In contrast, others did not show favourable results for blood loss and transfusion requirements when heparin titration was compared with an ACT-based protocol.^{152,161–163} Due to the lack of perioperative bleeding and transfusion rates as primary end points in these studies, larger multicentre RCTs are required to determine the added value of individual heparin management. Alternative methods of heparin management and anticoagulation monitoring during ECC consist of the determination of anti-factor Xa activity. The determination of thrombin levels could be promising, but it is not useful with high concentrations of heparin and is currently not available as a bedside clinical test.

Heparinization Algorithms. There is increasing interest in the application of heparinization algorithms to tailored heparin and protamine management in cardiac surgery. Application of these algorithms might contribute to improved anticoagulation strategies and PBM, especially in heparin-resistant patients, but RCTs evaluating the effectiveness of these models are currently lacking.

Measures to Prevent Heparin Rebound. Heparin rebound is the result of residual heparin blood concentrations in the postoperative phase. Although most of the literature on this topic originates from the 1980s to the 1990s, the recent literature can be divided into studies focusing on the presence of residual heparin in the blood following weaning from CPB or on the occurrence of postoperative bleeding associated with residual heparin. There is only 1 comparative study that showed

that protamine infusion (25 mg/h for 6 h) to neutralize residual heparin resulted in reduced 12-h blood loss compared with the control subjects (525 ± 322 vs 608 ± 385 ml, respectively; $P < 0.05$) but without affecting transfusion rates.¹⁶⁴ The study was, however, limited by the possibility of administering additional protamine to normalize ACT values to preheparin values, which occurred more frequently in the control group and might have enhanced postoperative bleeding.¹⁶⁴

In summary, a statement regarding optimal heparin anticoagulation management is limited by the lack of prospective and comparative studies. Heparin level-guided heparin management should, however, be considered over ACT-guided heparin management to reduce bleeding, especially in patients who are resistant to heparin.

5.3.2. Protamine

Background. The heparin–protamine complex leads to the dissociation of heparin from AT and restoration of the procoagulant properties of blood. Inadequate protamine dosing may, however, influence patient haemostasis and the risk of postoperative bleeding. The dose of protamine is usually based on the initial or total administered dose of heparin throughout the procedure. Protamine administration may be associated with immunological and inflammatory alterations and may induce hypotension, bradycardia, pulmonary vasoconstriction and allergies, which can, in most cases, be prevented by slow infusion rates, prophylactic use of ASA or an antihistaminic drug.

Description of the Evidence. In a small RCT, protamine dosing based on the initial heparin dose resulted in prolonged clotting times and microvascular bleeding compared with protamine dosing based on the measured heparin concentration following CPB.¹⁶⁵ In a second RCT, a protamine-to-heparin dosing ratio of 1.3 over the total heparin dose was associated with significantly more postoperative bleeding (615 ml, 95% CI 500–830 vs 470 ml, 95% CI 420–530; $P = 0.02$) compared with a dosing ratio of 0.8, whereas post-protamine ACT levels were comparable among the groups.¹⁵⁴ None of the patients showed signs of heparin rebound. A recent RCT in patients having CABG showed that a protamine-to-heparin dosing ratio < 0.6 was associated with significantly more blood loss within 12 h after surgery [420 ml (interquartile range 337–605 ml) vs 345 ml (interquartile range 230–482 ml); $P = 0.0041$], but no difference in PRBC transfusion was noted (2.3 units vs 2.7 units; $P = 0.83$) compared to patients subjected to a ratio > 0.8 .¹⁵² Ideally, protamine should be administered in a dose that matches actual heparin levels after the termination of ECC. It is advised not to exceed a protamine dose in a 1:1 ratio to the initial heparin bolus, because protamine overdosing might be associated with perioperative bleeding and enhanced transfusion requirements.

5.3.3. Antithrombin

Background. Patients resistant to heparin may have insufficiently low AT levels or dysfunctional AT and may require additional heparin dosing to reach the target ACT with a concomitant risk of residual heparin in the postoperative

phase. Alternatively, AT levels can be restored by recombinant AT supplementation before CPB.

Description of the Evidence. In a small RCT ($n = 20$), preoperative AT supplementation (50 U/kg) was not associated with reduced post-operative bleeding compared with the controls. At 6 and 12 h following surgery, median blood loss was similar between the groups ($P = 0.33$ and $P = 0.24$, respectively), with a higher variation in the AT group.¹⁶⁶ In heparin-resistant patients (defined as those demonstrating an inability to reach a target ACT of 480 s upon the administration of 300 + 100 IU/kg heparin), a single bolus of AT (75 U/kg) was associated with a reduced use of FFP. Administration of AT compared with placebo tended to increase 12-h postoperative bleeding (mean blood loss 1290 vs 756 ml; $P = 0.05$), without differences in transfusion rates between the groups.¹⁵⁵ In a second RCT, AT supplementation also resulted in higher bleeding rates 12 h after surgery compared with the controls (mean blood loss 450 vs 350 ml, respectively; $P = 0.011$).¹⁶⁷

In summary, AT supplementation is indicated in patients with AT deficiency to improve heparin sensitivity but should not be used prophylactically to reduce bleeding following CPB. FFP may be considered an alternative to AT supplementation in patients with AT deficiency to improve heparin sensitivity.

5.3.4. Heparin-Induced Thrombocytopenia Anticoagulation Management

Background. In patients with Type 2 heparin-induced thrombocytopenia (HIT), antibodies are generated against the antigen complex of heparin and platelet factor 4. This antigen-antibody complex may cause haemostatic and platelet activation and severe thromboembolism, which is diagnosed by highly specific platelet function tests or heparin-platelet factor 4 antibody enzyme-linked immunosorbent assays and platelet serotonin release assays.¹⁶⁸ Usually, HIT antibodies develop after 7–14 days of exposure to heparin with a half-life of 40–100 days. Newer generation ELISA tests are immunoglobulin G-specific, whereas older assays also measured immunoglobulin M and immunoglobulin A antibodies, resulting in a significant overdiagnosis of HIT.^{168–170} When HIT is diagnosed, heparin should be replaced by an alternative anticoagulant. The risks of alternative anticoagulation procedures, including increased bleeding, have to be balanced against potential HIT-associated complications.

Description of the Evidence. Direct thrombin inhibitors like bivalirudin or argatroban can be used as an alternative anticoagulant during CPB. Bivalirudin is the only agent that has been studied in prospective multicentre trials in patients with and without HIT undergoing on-pump and off-pump cardiac surgery.^{171–174} In a cardiac surgery trial in patients without HIT, safety data for bivalirudin were almost comparable to those of the competitor heparin-protamine.¹⁷³ Bivalirudin has a short elimination half-life (rv25 min) and no specific antidote. Due to its unique pharmacological characteristics, the surgical and perfusion strategies must be adjusted for the

target ACT, use of prime volume and cardioplegia to avoid the stagnation of blood in the CPB circuits.¹⁷⁵

Another strategy is to combine heparin with a potent shortacting antiplatelet agent, such as iloprost or tirofiban, to inhibit or attenuate the HIT reaction.^{176,177} However, when the effects of iloprost or tirofiban are eliminated, patients may again be at risk of developing HIT-associated complications. There are no sufficient data to support reduction of HIT antibodies via perioperative plasmapheresis.^{178,179}

In summary, in patients with HIT antibodies, in whom surgery cannot be postponed, bivalirudin anticoagulation may be considered when the bleeding risk is acceptable. The use of heparin in the pre- and postoperative periods should be avoided. If bivalirudin is deemed to be associated with a high risk of bleeding, heparin in combination with an antiplatelet agent may be considered.

5.4. Intravascular Volume

Recommendations	Class ^a	Level ^b	Ref ^c
The use of goal-directed haemodynamic therapy to reduce transfusions is not recommended.	III	C	
The use of modern low-molecular weight starches in priming and non-priming solutions to reduce bleeding and transfusions are not recommended.	III	C	
Limitation of haemodilution is recommended as part of a blood conservation strategy to reduce bleeding and transfusions.	I	B	180–182
Preoperative autologous blood donation in patients with high haemoglobin levels (> 110 g/l) may be considered to reduce postoperative transfusions.	IIb	B	183–185
Acute normovolaemic haemodilution may be considered to reduce postoperative transfusions.	IIb	B	186

^aClass of recommendation.

^bLevel of evidence.

^cReferences.

5.4.1. Goal-Directed Haemodynamic Therapy

Background. Goal-directed therapy is defined as the use of fluids, inotropes, vasoconstrictors and/or blood transfusions to target systemic haemodynamic goals, such as blood pressure or cardiac output, which improve the perfusion and oxygenation of tissues.

Description of the Evidence. An RCT combined with a systematic review¹⁸⁷ was designed to investigate the effect of goal-directed therapy on outcomes in high-risk cardiac surgical patients (EuroSCORE > 6 and/or with recent MI < 14 days).

Although patients in the goal-directed therapy arm received more intravenous fluids compared with the controls, there were no differences in intraoperative PRBC transfusion volumes between the groups [0 (0) units vs 1 (1.6) units, respectively; $P = 0.32$].¹⁸⁷ There are currently no other studies available in this area that focus on bleeding and blood transfusions.

Low blood pressure is, among other interventions, treated by vasopressors, but the rise in blood pressure might contribute to enhanced blood loss. On the other hand, maintaining the mean arterial pressure using a vasoconstrictor mitigates the haemodilution associated with anaesthesia and CPB.¹⁸⁸ One RCT showed that goal-directed therapy using fluids, inotropes and vasopressor therapy did not affect the secondary end point transfusion requirements, despite there being more vasopressor therapy than in the control group.¹⁸⁹ Given the availability of limited data, it is not possible to reach evidence-based conclusions on the benefits of goal-directed fluid therapy in this setting.

5.4.2. Use of Crystalloids and Colloids

Background. There is no consensus regarding optimal fluid strategies in cardiac surgery, but it has been suggested that the use of hydroxyethyl starches (HES) may be associated with increased perioperative blood loss. Most of the available RCTs are limited by sample size and focus on the effect of different priming solutions on patient haemostasis, with bleeding or transfusion requirements as a secondary end point.

Description of the Evidence

Priming Solutions. In studies addressing the impact of the type of priming solution on postoperative bleeding and blood product transfusion as secondary end points, a comparison of 6% HES 130/0.4 (mean molecular weight 130 kDa; degree of substitution) with 6% HES 400/0.7 prime with gelatine prime,¹⁹⁰ albumin prime^{191,192} or Ringer's acetate¹⁹³ was made. There was no effect of 6% HES 130/0.4 on blood loss or transfusion requirements. Two studies showed a reduction in postoperative blood loss when albumin was added to an HES 130/0.4 prime¹⁹⁴ or compared to 6% HES 200/0.5 with 3.5% gelatin.¹⁹⁵ However, a review of prospective studies that compared different priming solutions with blood loss and/or transfusion requirements as primary end points reported no difference in postoperative blood loss when different HES solutions were evaluated.^{196–199}

Volume Therapy. Two studies investigated the impact of different intraoperative volume therapy strategies on postoperative bleeding and/or transfusion as the primary end point. A comparison of 1 litre of 6% HES with 5% albumin during off-pump CABG procedures was associated with significantly more 12-h chest drainage volume (mean blood loss 732 vs 563 ml; $P < 0.001$).²⁰⁰ In contrast, 6% HES 130/0.4 vs 3.5% gelatine showed no differences in total blood loss (mean blood loss 19.4 vs 19.2 ml/kg).²⁰¹ In a 3-armed study, there was no difference in 24-h blood loss when 5% albumin or Ringer's lactate was compared with 6% HES 130/0.4 (mean blood loss 835 vs 670 vs 700 ml, respectively; $P = 0.085$). However, the transfusion rates of PRBC were higher in the albumin and

HES groups compared with that of Ringer's lactate (58% vs 61% vs 24%, respectively; $P = 0.0013$), with the highest creatinine levels reported for the HES group.²⁰²

In summary, in studies designed to show a difference in blood loss or transfusion requirements, distinct priming solutions or perioperative volume strategies revealed comparable results. However, due to the suggestion that the use of HES may be associated with kidney failure and increased mortality rates, starches are decreasingly used as volume therapy during cardiac surgery. It is therefore not recommended to use modern low-molecular-weight starches in priming and non-priming solutions to reduce bleeding and transfusions.

5.4.3. Haemodilution and Cardioplegia

Background. Haemodilution is advantageous to a certain degree, because it lowers blood viscosity and may improve microcirculatory perfusion. However, haemodilution also leads to a reduced concentration of coagulation factors and blood cells during and following ECC, which may have implications for postoperative haemostasis and organ function.

Description of the Evidence. Publications from large database studies have shown that low haematocrit values are associated with increased morbidity¹⁸² and in-hospital deaths.²⁰³ One meta-analysis showed that the use of minimized circuits reduces haemodilution, postoperative bleeding and transfusion requirements.¹⁰⁰ Two studies focused on haemodilution independent of the use of MiECC systems by implementing a multimodality blood conservation scheme or restricting intravenous fluids during surgery.^{180,181} In both studies, a marked reduction in the number of transfusions with increased haematocrit values was achieved.^{180,181}

In conclusion, limitation of haemodilution is recommended as part of a blood conservation strategy to reduce bleeding and transfusions.

5.4.4. Predonation of Blood

Background. Preoperative autologous blood donation in the hours, days or weeks before surgery may reduce the number of allogeneic blood transfusions. Preoperative blood donation is limited to patients with relatively high Hb and haematocrit levels and with a relatively large body surface area without coagulation abnormalities.

Description of the Evidence. A large retrospective study showed that autologous blood donation in elective CABG or valve surgery was associated with a lower occurrence of allogeneic blood transfusion and cost-effectiveness.¹⁸⁴ However, the study lacked statistical adjustment for confounding factors.¹⁸⁴ In a matched-pair analysis with 432 patients, it was shown that preoperative autologous blood donation was associated with a reduction in the transfusion rate of PRBC from 55% to 32% and a 50% reduction in the administration of FFP and platelet concentrate (PLTC).¹⁸³ A nested case-control study (corrected for confounding factors) further revealed that preoperative autologous blood donation reduced PRBC transfusions by 18.3% in valve surgery.²⁰⁴ This study was,

however, limited by the lack of preoperative Hb and haematocrit data.²⁰⁴

Based on the current evidence, we suggest that preoperative autologous blood donation in patients without severe aortic stenosis, Canadian Cardiovascular Society (CCS) Grade 3–4 angina or ACS < 4 weeks and with high Hb levels (> 110 g/l) who are having elective surgery may be considered to reduce the number of postoperative transfusions.

5.4.5. Acute Normovolaemic Haemodilution

Background. During acute normovolaemic haemodilution (ANH), the patient donates blood just before the onset of surgery that is replaced by an equal amount of volume to maintain normovolaemia. The donated blood is retransfused after CPB, but this practice comes at the cost of lower haematocrit values during surgery.

Description of the Evidence. A recent systematic review and meta-analysis including 29 RCTs investigated the effect of ANH using a variation of colloid, albumin and crystalloid volume replacement on allogeneic PRBC transfusion requirements in adult and paediatric cardiac surgical procedures.¹⁸⁶ The majority of the studies included had no predefined transfusion protocols. In the 21 studies with allogeneic blood transfusion as a predefined end point, patients subjected to ANH received fewer PRBCs (-0.79 units; 95% CI -1.25 to -0.34; $P = 0.001$), albeit with a considerable variation in the effect size between studies.¹⁸⁶ Moreover, most of the studies with a large benefit of ANH on PRBC transfusion were published before 2001. In the studies reported after 2001, only half of the studies showed a benefit of ANH on PRBC transfusion. ANH was associated with a significant, but clinically irrelevant, reduction in postoperative bleeding (-0.64 ml; 95% CI -0.97 to -0.31; $P < 0.0001$).

In summary, ANH may have a beneficial effect on postoperative transfusion rates, but this conclusion should be viewed in light of the older studies included in this meta-analysis.

6. Coagulation and Transfusion

Postoperative diagnosis and guidance of patient haemostasis and treatment of haemostatic abnormalities by specific procoagulant interventions or blood transfusion products should be considered as closing the circle of PBM in cardiac surgery.

6.1. Procoagulant Interventions

Recommendations	Class ^a	Level ^b	Ref ^c
Antifibrinolytic therapy (TXA, aprotinin and EACA) is recommended to reduce bleeding and transfusions of blood products and reoperations for bleeding (TXA and aprotinin).	I	A	205–207

The prophylactic use of FFP to reduce bleeding is not recommended.	III	B	208–210
The use of PCC or FFP may be considered to reverse the action of VKAs.	IIb	B	211,212
In patients with factor XIII activity < 70% after CPB, the administration of factor XIII may be considered to reduce bleeding.	IIb	B	213,214
Prophylactic fibrinogen administration is not recommended.	III	B	215
In the bleeding patient with a lowfibrinogen level (< 1.5 g/l), fibrinogen substitution may be considered to reduce postoperative bleeding and transfusions.	IIb	B	216–218
In patients where bleeding is related to coagulation factor deficiency, PCC or FFP administration should be considered to reduce bleeding and transfusions.	IIa	B	219,220
The prophylactic use of DDAVP to reduce bleeding is not recommended.	III	B	221,222
In bleeding patients with platelet dysfunction on the basis of an inherited or acquired bleeding disorder, the use of DDAVP should be considered to reduce bleeding and the requirement for transfusions.	IIa	C	
The prophylactic use of rFVIIa to prevent bleeding is not recommended.	III	B	223
In patients with refractory, non-surgical bleeding, off-label use of rFVIIa may be considered to reduce bleeding.	IIb	B	224

Bleeding is defined as persistent, non-surgical microvascular blood loss.

^aClass of recommendation.

^bLevel of evidence.

^cReferences.

CPB: cardiopulmonary bypass; DDAVP: desmopressin; EACA: ϵ -amino-caproic acid; FFP: fresh-frozen plasma; PCC: prothrombin complex concentrate; rFVIIa: recombinant activated factor VII; TXA: tranexamic acid; VKAs: vitamin K antagonists.

6.1.1. Antifibrinolytics

Background. Antifibrinolytic therapy is widely used to reduce bleeding, transfusions of blood products and reoperations for bleeding. Three antifibrinolytics are used to reduce reoperation for bleeding and blood product transfusion in cardiac surgery: TXA, aprotinin and EACA. Although aprotinin was withdrawn from the market in 2007, in some countries, the drug remains available through a limited access plan.

Tranexamic Acid. The Aspirin and Tranexamic Acid for Coronary Artery Surgery (ATACAS) RCT compared TXA with placebo in patients having CABG surgery and

demonstrated a reduction in the risk for reoperation due to major haemorrhage (RR 0.36, 95% CI 0.21–0.62; $P < 0.001$) and in the need for the transfusion of any blood products (37.9% vs 54.7%; $P < 0.001$).²⁰⁵ This result confirms the findings of previous meta-analyses.^{206,225,226} In a systematic review and cumulative meta-analysis, it was shown that TXA reduced the pooled risk ratio for blood transfusion in patients having cardiac surgery compared with the controls (RR 0.65, 95% CI 0.60–0.70; $P < 0.001$).²²⁶ The most significant reported side effect of TXA was convulsive seizure.^{227,228} The findings from the ATACAS trial showed that low-dose (50 mg/kg) versus high-dose (100 mg/kg) TXA was not safer in terms of seizures (0.7 vs 0.6%, respectively), but the higher dose significantly reduced blood loss ($P = 0.026$) and PRBC transfusion ($P = 0.017$).²⁰⁵

Aprotinin. The use of aprotinin in patients having cardiac surgery has been studied extensively. Several RCTs and meta-analyses demonstrated its efficacy in reducing the risk of reoperation for bleeding (RR 0.46, 95% CI 0.34–0.63; $P < 0.001$) and the need for blood product transfusion (RR 0.68, 95% CI 0.63–0.73; $P < 0.001$).²⁰⁶ It was suggested that aprotinin was a more effective agent than lysine analogues in reducing allogeneic blood exposure, but the differences between the antifibrinolytics were small (RR 0.86, 95% CI 0.76–0.96; $P = 0.0083$).²⁰⁶ Moreover, most of the studies were restricted to patients having CABG.²⁰⁷

Data from observational studies^{229,230} indicated an increased risk of death associated with aprotinin. In these studies, aprotinin was preferentially administered to patients with more risk factors for increased mortality rates at the baseline compared with the control groups. Several observational studies did not report an association between aprotinin and safety outcomes.^{231–233} The Blood Conservation Using Antifibrinolytics in a Randomized Trial (BART),²³⁴ which was not designed to look at safety, reported higher mortality rates for patients treated with aprotinin compared to those treated with TXA or EACA, which led to the regulatory marketing suspension of aprotinin. Subsequent recognition of the significant limitations of the BART trial, including crossover of patients in the trial arms,²³⁵ and a network meta-analysis showing no adverse effect of aprotinin compared with placebo and other agents,²³² led to the lifting of the marketing restriction in Canada and by the European Medicine Agency in 2012 in Europe.^{236,237}

ϵ -Aminocaproic Acid. In comparison with TXA, limited data are available to support the use of EACA in cardiac surgery.²⁰⁶ A meta-analysis of observational studies showed that the use of EACA reduced the risk for blood product transfusions (RR 0.70, 95% CI 0.52–0.93; $P = 0.015$) but did not decrease the risk of re-exploration for bleeding (RR 0.35, 95% CI 0.11–1.17; $P = 0.087$).

In summary, safety studies should be designed to define the dose–response relationship of the adverse effects observed following the administration of aprotinin and TXA, given the positive impact these agents have on bleeding.

6.1.2. Fresh-Frozen Plasma

Background. Fresh frozen plasma is obtained by separating the plasma, including coagulation factors and other proteins

and soluble constituents of the blood, from particular elements in donated blood. An increasing number of countries use pooled plasma. Pooled plasma contains plasma from multiple donors and is inactivated for viruses with a lower risk of transfusion-induced lung injury.

Description of the Evidence. The largest body of evidence is gathered on the prophylactic administration of FFP to patients without a diagnosed coagulopathy and is summarized in a Cochrane review²⁰⁸ and 3 other systematic reviews.^{209,210,238} The RCTs included were limited by small sample sizes and divergent doses of FFP. There was no difference in blood loss or allogeneic blood transfusion in patients undergoing cardiac surgery who received FFP compared with the controls.^{209,210,238} It was also shown that the therapeutic use of FFP in patients with established coagulopathy or for reversal of oral anticoagulation has no effect on 24-h blood loss.^{209,238}

In summary, FFP might be used to reverse the action of oral anticoagulation or in the case of persistent perioperative bleeding, but there is no evidence that prophylactic or therapeutic FFP transfusions reduce blood loss after cardiac surgery.

6.1.3. Factor XIII

Background. Factor XIII (FXIII) is the terminal enzyme in the coagulation cascade and is necessary for cross-linking fibrin monomers to form a stable, strong fibrin clot. Low FXIII levels have been reported after cardiac surgery and are associated with increased bleeding.^{213,239,240} Specific immunological assays are needed to determine functional FXIII activity. Detection of FXIII deficiency by TEM has been described,²⁴¹ but there is currently no clinically available point-of-care (POC) test.

Description of the Evidence. Three RCTs ($n = 527$) evaluated the effect of the administration of FXIII after the administration of protamine.^{213,242,243} The results showed no differences in postoperative bleeding volumes and transfusion rates with any FXIII dosage regimens. There was no difference in adverse events, including thromboembolic events and death, between the intervention and the control groups. However, in patients with a postoperative FXIII level 70%, the bleeding volume was higher compared to patients with FXIII levels $> 70\%$ ²¹³; therefore, the latter patients might be the target population.

In summary, there is no evidence that administration of FXIII might be beneficial regarding the reduction of postoperative bleeding volumes and transfusion rates in patients with normal concentrations of FXIII ($> 70\%$).

6.1.4. Fibrinogen Concentrates

Background. Fibrinogen is the first factor to be depleted during significant bleeding and haemodilution.^{220,244} Preoperative fibrinogen levels < 1.5 g/L²⁵ and hypofibrinogenemia after cardiac surgery^{242,245} have been associated with increased postoperative bleeding. Fibrinogen supplementation

has, therefore, been advocated as one of the primary haemostatic aims in patients who bleed after cardiac surgery.²⁴⁶

Description of the Evidence

Pre- and Postoperative Fibrinogen Substitution. In an RCT including 48 patients, preoperative fibrinogen substitution slightly increased intra- and postoperative fibrinogen concentrations but did not reduce postoperative bleeding in patients who had CABG (mean blood loss 650 vs 730 ml, $P = 0.29$).²⁴⁷ Postoperative fibrinogen administration was tested in 4 studies, with contrasting results. Two studies^{248,249} that included 167 patients undergoing complex cardiac surgery showed reduced requirements for allogeneic blood products in the intervention group. In both studies, fibrinogen concentrate was administered, the goal being a maximal clot firmness of 22 mm in FIBTEM (ROTEM, Basel, Switzerland). These findings are in agreement with a Cochrane systematic review published in 2013, which showed a reduced relative risk of blood transfusion after fibrinogen supplementation (RR 0.47, 95% CI 0.31–0.72).²¹⁶ However, 2 more recent RCTs in bleeding patients after complex cardiac surgery could not confirm the previous findings.^{215,250} Indeed, the largest RCT of 152 bleeding patients without hypofibrinogenaemia undergoing elective aortic surgery, randomly assigned to fibrinogen concentrate or placebo, found that patients treated with a fibrinogen concentrate had a significantly higher rate of any allogeneic blood product use compared with placebo (median units 5.0 vs 3.0, $P = 0.026$).²⁵⁰ A *post hoc* analysis from an RCT showed that substitution with fibrinogen concentrate to more than 2.87 g/l (or maximal clot firmness of 14 mm in FIBTEM) could not further reduce bleeding volume.²⁵¹ Data from a Cochrane review including 124 patients from 3 RCTs²¹⁶ and a propensity score analysis in 380 patients undergoing cardiac surgery²⁵² suggested no increased risk of thromboembolic events or death after fibrinogen substitution. However, additional safety data from larger RCTs, which also addressed different fibrinogen target concentrations, are necessary to confirm this assumption.

In summary, prophylactic fibrinogen administration is not recommended for reducing postoperative bleeding and transfusion risks. However, in patients with a low-fibrinogen level and signs of persistent microvascular bleeding, fibrinogen substitution may be considered to reduce the requirement for transfusions.

6.1.5. Prothrombin Complex Concentrate

Background. PCCs are lyophilized, human plasma-derived vitamin K-dependent coagulation factors. In Europe, only 4 factor concentrates are available, containing coagulation factors II, VII, IX and X. Although the concentration of factor IX is standardized, the other factors might vary among the different available products. Also, PCC might contain various amounts of AT; proteins C, S and Z and heparin.^{246,253}

Description of the Evidence. In patients with extremely high INR values (> 4 –5) taking VKAs for the long term who need urgent or emergent surgery, the use of PCCs might be

advantageous compared with FFP due to the more rapid reversal of the anticoagulant effect. A Cochrane review, however, showed no evidence of a difference between PCC and FFP in terms of postoperative bleeding and transfusion risks.²¹² In 40 patients with an INR > 2.1 , the perioperative administration of PCC reversed anticoagulation safely; reversal was faster and postoperative bleeding volume was lower compared with the administration of FFP.²¹¹

Three retrospective cohort studies in 402 patients showed that PCC was associated with fewer re-explorations for bleeding, less blood loss and fewer transfusions of PRBCs compared with FFP.^{219,254,255} The use of PCCs in patients with bleeding after CPB might be associated with an increased risk of thromboembolic events and acute kidney injury.²⁵⁵ Recommendations for dosing are not established, and administration might be tailored to the individual patient based on laboratory and clinical variables, including POC coagulation testing.^{246,253}

To summarize, in patients in whom bleeding is related to coagulation factor deficiency, PCC or FFP administration should be used to reduce bleeding and transfusions. PCC may be preferred over FFP when rapid normalization of coagulation factors is needed.

6.1.6. Desmopressin

Background. Desmopressin (DDAVP) is a haemostatic agent that was developed to prevent and treat bleeding in patients with mild haemophilia (5–50% of the normal concentration of clotting factors VIII and IX) and patients with von Willebrand's deficiency, e.g. due to aortic stenosis. DDAVP stimulates the release of von Willebrand factor from endothelial cells. DDAVP is sometimes administered to treat postoperative bleeding in patients undergoing cardiac surgery with impaired platelet function.

Description of the Evidence. The effects of DDAVP on bleeding in patients having cardiac surgery are summarized in a Cochrane systematic review of 29 RCTs²⁵⁶ and 2 meta-analyses.^{222,257} Summing up the evidence, the prophylactic use of DDAVP did not reduce blood loss (weighted mean difference 241 ml, 95% CI -387.55 to -96.01 ml) or the risk of blood transfusions (OR 0.96, 95% CI 0.87–1.06). Recently published meta-analyses concluded that DDAVP has a small effect on the volume of blood loss and allogeneic blood product transfusions.^{222,257} Furthermore, the effect of DDAVP may be more profound in subgroups, e.g. for patients on platelet inhibitors, patients with reduced platelet function and patients exposed to prolonged CPB times.²²² There was no increased risk for MI (RR 1.38, 95% CI 0.77–2.50), stroke (RR 2.40, 95% CI 0.68–8.43) or any thromboembolic complications (RR 1.46, 95% CI 0.64–3.35) after the administration of DDAVP.^{256,257} However, the risk of hypotension during administration of DDAVP that required fluids and/or vasoactive drugs was significantly increased (RR 2.81, 95% CI 1.50–5.27).²⁵⁶

In summary, prophylactic use of DDAVP is not recommended, but its administration might be beneficial in patients

who are bleeding who have inherited or acquired bleeding disorders or platelet dysfunction.²⁵⁸

6.1.7. Recombinant Factor VIIa

Background. Recombinant factor VIIa (rFVIIa) is used in the treatment and prevention of bleeding in patients with inherited bleeding disorders. The observed haemostatic properties of this agent led to its off-label use in life-threatening bleeding, but nowadays its use is considered to be a last resort in the event of uncontrollable blood loss.

Description of the Evidence. A 2012 Cochrane review analysed 29 RCTs with respect to the prophylactic (1361 patients) and therapeutic use (2929 patients) of rFVIIa in patients with or without haemophilia.²²⁴ Prophylactic rFVIIa use just failed to reach significance to reduce transfusion rates (RR 0.85, 95% CI 0.72–1.01) compared with placebo, but this result was associated with an increased risk of thromboembolic adverse events (RR 1.35, 95% CI 0.82–2.25). There was no benefit of the therapeutic use of rFVIIa compared with the use of placebo with respect to transfusion rates, but the administration of rFVIIa was associated with reduced mortality rates (RR 0.91, 95% CI 0.78–1.06) at the cost of more major thromboembolic events (RR 1.14, 95% CI 0.89–1.47).

In summary, prophylactic use of rFVIIa cannot be recommended in cardiac surgery; its therapeutic use should only be considered in patients with uncontrollable bleeding that cannot be managed by other procoagulant interventions.

6.2. Transfusion Strategies

Recommendations	Class ^a	Level ^b	Ref ^c
Implementation of a PBM protocol for the bleeding patient is recommended.	I	C	
The use of PRBCs of all ages is recommended, because the storage time of the PRBCs does not affect the outcomes.	I	A	259,260
The use of leucocyte-depleted PRBCs is recommended to reduce infectious complications.	I	B	261
Pooled solvent detergent FFP may be preferred to standard FFP to reduce the risk of TRALI.	IIb	B	262
Perioperative treatment algorithms for the bleeding patient based on viscoelastic POC tests should be considered to reduce the number of transfusions.	IIa	B	263–265
It is recommended that one transfuse PRBCs on the basis of the clinical condition of the patient rather than on a fixed haemoglobin threshold.	I	B	266,267
A haematocrit of 21–24% may be considered during CPB when an	IIb	B	268

adequate DO₂ (> 273 ml O₂/min/m²) level is maintained.
Platelet concentrate should be transfused in bleeding patients with a platelet count below 50 (10⁹/l) or patients on antiplatelet therapy with bleeding complications.

^aClass of recommendation.
^bLevel of evidence.
^cReferences.
CPB: cardiopulmonary bypass; FFP: fresh-frozen plasma; PBM: patient blood management; POC: point of care; PRBCs: packed red blood cells; TRALI: transfusion-related acute lung injury.

6.2.1. Quality of Blood Products

Background. Stored PRBCs or FFP reveal comparable efficacy compared to fresh products but differ in terms of the risk of the transmission of bacterial or viral infections,^{269,270} transfusion-related acute lung injury (TRALI)^{271,272} and transfusion-related immune modulation (TRIM).^{273–275} To avoid TRALI, FFP and PLTC must be collected from men, women who have not been pregnant and women who have tested negative for human leucocyte antigen antibodies to reduce the risk of TRALI.²⁷¹ The effects of TRIM are assumed to be caused by allogeneic mononuclear cells, white blood cell-derived soluble mediators and human leucocyte antigen peptides, transfused mainly via PRBCs.²⁷³ TRIM has been associated with increased rates of mortality and morbidity in cardiac surgery and an increased risk of developing infections.^{274,275}

Description of the Evidence. Leucocyte reduction of PRBC is the standard practise in Europe and preserves the quality of PRBC during storage and influences favourably on morbidity.²⁶¹ In an RCT, patients having cardiac surgery either received buffy coat-depleted or leucocyte-depleted erythrocytes.²⁷⁶ Although no differences in mortality rates could be shown, patients receiving leucocyte-depleted PRBCs showed a reduced infection rate compared with the controls (21.6% vs 31.6%; OR 1.64, 95% CI 1.08–2.49).²⁷⁶ The implementation of new additive solutions further improved PRBC quality during storage.

In the Red-Cell Storage Duration Study (RECESS), 1098 cardiac surgical patients were randomized to receive exclusively ‘new’ (≤ 10 days) or ‘old’ (≥ 21 days) leucocyte-depleted PRBCs.²⁵⁹

There were no differences in multiple-organ dysfunction and mortality rates between either arm. These results are in line with the results of a large Swedish registry that compared the outcomes of 47 071 cardiac surgery patients from 1997 to 2012 who were exclusively transfused with PRBC stored for < 14 days (36.6%), 14–27 days (26.8%), 28–42 days (8.9%) or of mixed age (27.8%).²⁷⁷ No differences were noted in terms of 30-day-, 2-year- and 10-year mortality rates or in relation to 30-day organ dysfunction or serious infection.²⁷⁷ A current meta-analysis that included the Red-Cell Storage Duration Study²⁹¹ and the large Age of BLOOD Evaluation

(ABLE) trial²⁶⁰ showed that red blood cell storage time does not affect mortality rates, adverse events or nosocomial infections.²⁷⁸

PLTCs can be produced as single-donor apheresis PLTC or as pooled (4–5 donors) whole-blood-derived PLTC. Although the risk for TRALI appears to be comparable, an increased risk of transmitting infections is associated with whole-blood-derived PLTC transfusions.²⁷⁹ One retrospective study in 3272 patients who received single-donor PLTC aged 2–5 days showed no association between PLTC storage age and short-term outcome, survival or postoperative infections.²⁸⁰ Solvent–detergent treatment and pooling of FFP (SD-FFP) are performed to inactivate capsulated lipid-enveloped viruses and dilute human leucocyte antigen antibodies to reduce TRALI. In a haemovigilance report from Norway (1998–2003), where SD-FFP is exclusively used, no case of TRALI has been associated with SD-FFP transfusion, whereas 3 other European countries have reported incidences of 1.6–8.8 when transfusing FFP.²⁶²

In conclusion, we recommend the use of PRBCs of all ages, because the storage time of the PRBCs does not affect outcomes. In an attempt to reduce infectious complications, the use of leucocyte-depleted PRBCs is recommended; in contrast, the pooled SD-FFP may be preferred over standard FFP to reduce the risk for TRALI.

6.2.2. Algorithm-Guided Therapy of Perioperative Bleeding Background. The introduction of POC tests—in particular, TEG and TEM—has led to the development of specific

algorithms to diagnose and, consequently, treat theragnostic perioperative bleeding in cardiac surgery. The first algorithms were based on TEG, and their effectiveness was tested in a small series using historical controlled series, fibrinogen concentration measurements using the Clauss method or a viscoelastic fibrinogen test.^{251,281} Figure 3 provides an overview of possible monitoring options throughout the perioperative period.

Description of the Evidence. A meta-analysis with 5223 patients shows that exposure to any allogeneic blood product was lower in the POC group (49.6%) than in the conventional group (60.2%; OR 0.63, 95% CI 0.56–0.71; $P < 0.0001$). This finding was confirmed when we considered RCTs only (OR 0.37, 95% CI 0.21–0.68; $P = 0.0018$). The most relevant decrease in transfusion rate was related to FFP (OR 0.31, 95% CI 0.13–0.74), followed by platelet transfusion (OR 0.62, 95% CI 0.42–0.92) and PRBCs (OR 0.63, CI 0.50–0.78).²⁶³ Data on chest tube drainage in a subpopulation of 400 patients revealed no differences. Data on the surgical re-exploration rate favoured the POC group versus the conventional group (OR 0.56, 95% CI 0.45–0.71; $P < 0.00001$).²⁶³ However, a recently published meta-analysis could not show any benefit of algorithm-based therapy on surgical reoperation rates.²⁶⁴ There was a reduction of thromboembolic events in the POC-guided group versus the conventional group (1.3% vs 2.9%; OR 0.44, 95% CI 0.28–0.70; $P = 0.0005$). No differences were found with regard to mortality rates (OR 0.92, 95% CI 0.74–1.16; $P = 0.45$).²⁶³ The majority of the studies considered

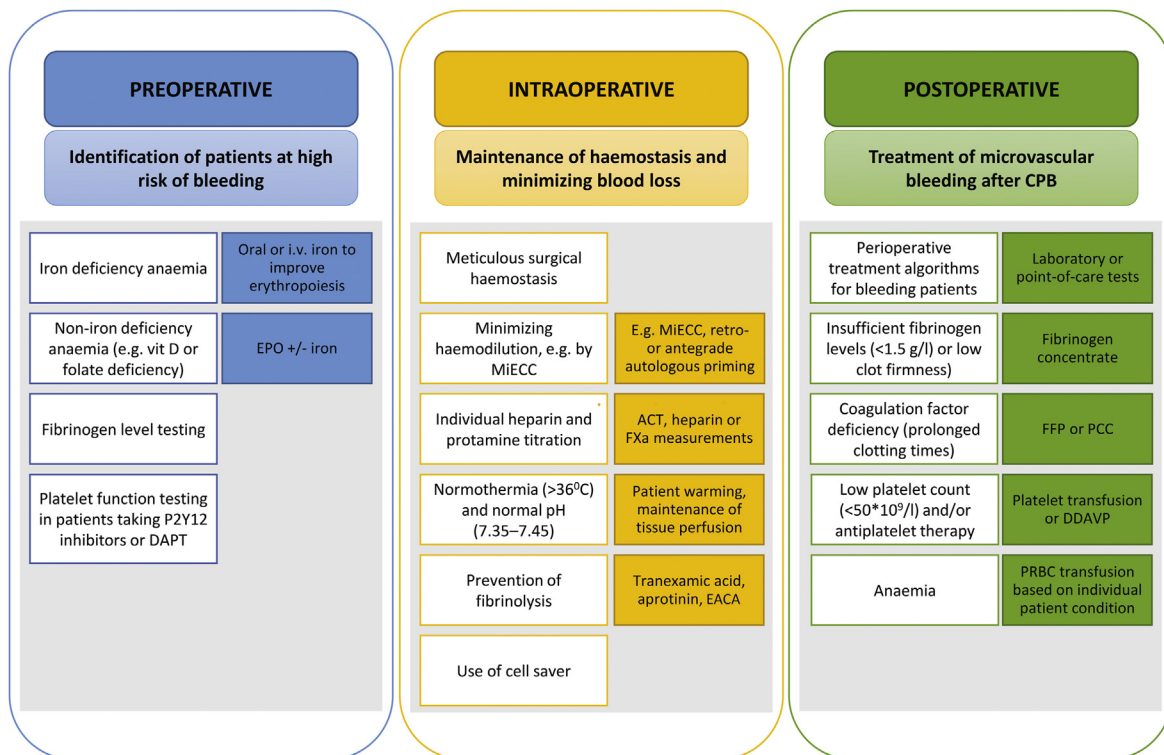


Figure 3. Haemostatic monitoring throughout the perioperative period and possible treatment modalities. ACT: activated clotting time; CPB: cardiopulmonary bypass; DAPT: dual antiplatelet therapy; DDAVP: desmopressin; EACA: ε-aminocaproic acid; EPO: erythropoietin; FFP: fresh-frozen plasma; i.v.: intravenous; MiECC: minimally invasive extracorporeal circulation circuit; PCC: prothrombin complex concentrate; PRBC: packed red blood cells.

algorithms based on viscoelastic tests, whereas a minority reported data inclusive of platelet function tests.²⁶³

In 2016, an updated comprehensive Cochrane report that primarily included cardiac surgery reports assessed the benefits and harms of TEG- and TEM-guided transfusions in adults and children with bleeding.²⁸² The use of TEG- and TEM-guided management was associated with a reduction in the proportion of patients receiving PRBC transfusions (RR 0.86, 95% CI 0.79–0.94), FFP (RR 0.57, 95% CI 0.33–0.96) and platelets (RR 0.73, 95% CI 0.60–0.88), whereas there was no significant difference in the rate of surgical reoperation or bleeding events and massive transfusions.²⁸² In 2015, a Dutch/British health technology assessment report confirmed the results of the Cochrane analysis and showed that viscoelastic testing was cost-effective and more efficient than standard laboratory testing.²⁸³ An additional meta-analysis of TEG-/TEM-based coagulation management included 9 RCTs and 8 observational studies comprising 8332 patients.²⁶³ The authors found that TEG-/TEM-based management reduced the odds for transfusion requirements (OR 0.63, 95% CI 0.56–0.71) and re-explorations for bleeding (OR 0.56, 95% CI 0.45–0.71) but made no difference in terms of mortality rate, stroke or chest drain output volume.²⁶³ In a subsequent RCT, 7402 patients were randomized to POC haemostatic testing with an integrated transfusion algorithm or standard of care.²⁶⁵ POC haemostatic testing reduced the rates of PRBC transfusions (adjusted RR 0.91, 95% CI 0.85–0.98), platelet transfusions (RR 0.77, 95% CI 0.68–0.87) and major bleeding (RR 0.83, 95% CI 0.72–0.94) without affecting complication rates.²⁶⁵

In summary, the evidence supports the use of perioperative POC testing in patients having cardiac surgery to reduce transfusion requirements. Furthermore, it should be noted that most of the published studies included only viscoelastic coagulation tests, although a specific platelet function test was included in the algorithm in 2 of the more recent studies.^{265,284} Cut-off levels for non-acceptable intraoperative and postoperative platelet function with the different devices and the subsequent interventions are yet to be determined.

6.2.3. Transfusion Triggers for Packed Red Blood Cells and Platelet Concentrate

Background. The transfusion threshold is usually defined as the critical haematocrit/Hb value, red cell volume or platelet count/function when patients are transfused with PRBCs or PLTC. However, dynamic strategies also take tissue perfusion/oxygenation into account.^{285,286} Data of RCTs in cardiac surgery are hard to compare, because transfusion thresholds of PRBC transfusion and the quality of blood products vary among studies.

Description of the Evidence. A recent meta-analysis compared restrictive (Hb 70–80 g/l) and liberal (Hb 90–100 g/l) transfusion strategies using data from 6 RCTs with 3352 patients, showing a 30% reduction in the number of deaths in the restrictive regimen group (OR 0.70, 95% CI 0.49–1.02; $P = 0.060$).²⁸⁷ The Brazilian Transfusion Requirements After Cardiac Surgery

(TRACS) trial compared a liberal transfusion threshold (haematocrit $\geq 30\%$) with a restrictive threshold (haematocrit $\geq 24\%$) in 502 patients having CABG and/or valve surgery.²⁶⁷ The per protocol analysis showed that more patients in the liberal group received red blood cell transfusions (78% vs 48%; $P < 0.001$) without a difference in the composite end point of 30-day mortality or severe morbidity.²⁶⁷ A subanalysis of a small sample of patients older than 60 years ($n = 260$) showed no difference in the composite end point between the groups, albeit with more cardiogenic shock in the restrictive transfusion threshold group (12.8% vs 5.2%, $P = 0.031$).²⁸⁸

The UK Transfusion Indication Threshold Reduction (TITRe2) trial compared a liberal (Hb < 90 g/l) with a restrictive threshold (Hb < 75 g/l) for leucocyte-depleted red blood cells in 2000 patients undergoing CABG and/or valve or aortic surgery.²⁶⁶ Fewer patients in the restrictive group received red blood cells (63.7% vs 94.9%; RR 0.58, 95% CI 0.54–0.62; $P < 0.001$), whereas the 90-day mortality rate was doubled in the restrictive group (4.2% vs 2.6%; OR 1.64, 95% CI 1.00–2.67; $P = 0.045$). The TITRe2 study was criticized regarding transfusion thresholds and confounders, which might be overcome in the ongoing Transfusion Requirements in Cardiac Surgery (TRICS III) study.

A small RCT conducted with patients undergoing CABG with normothermic CPB and not requiring transfusions compared the effect of a haematocrit of 20% ($n = 28$) with that of a haematocrit of 25% ($n = 26$). No differences in oxygen delivery or consumption, blood lactate levels or clinical outcome were found.²⁶⁸ In a single-centre investigation in which surgery was restricted to CABG ($n = 3003$), a nadir haematocrit $< 28\%$ during CPB was associated with an increase in severe morbidity, whereas lowering the nadir haematocrit to 24% was possible when a goal-directed perfusion technique was established, which included maintenance of sufficient oxygen delivery.¹⁸² These findings should highlight the complex interaction of haematocrit, temperature, oxygen supply and tissue perfusion for tissue oxygenation.

There are no studies available in which a certain platelet count or value in a platelet function assay served as a threshold for PLTC transfusion and could be identified as the key effector to stop/reduce microvascular bleeding caused by inadequate platelet count/function.²⁸⁹ However, in recent American Association of Blood Bank guidelines, a platelet count of 50 ($10^9/l$) was recommended as a therapeutic target for the bleeding patient undergoing major surgery.²⁹⁰

In summary, recent studies seem to be in favour of liberal PRBC transfusions. However, from an expert opinion, we would like to underline that the clinical condition of the patient and the optimization of the balance between oxygen delivery and extraction in the tissues are the most important aspects to be considered rather than a fixed threshold of the level of Hb. The acceptable haematocrit levels during CPB depend on the risk profile of the patient and his or her ability to maintain adequate tissue perfusion and oxygenation. In bleeding patients with a platelet count below 50 ($10^9/l$) or patients on antiplatelet therapy, PLTC may be transfused.

7. Anticoagulation Management During Extracorporeal Life Support

Recommendations	Class ^a	Level ^b	Ref ^c
Bivalirudin and argatroban may be considered as alternatives for heparin anticoagulation for prolonged ECLS/ECMO therapy to prevent HIT type 2.	IIb	C	

^aClass of recommendation.

^bLevel of evidence.

^cReferences.

ECLS: extracorporeal life support; ECMO: extracorporeal membrane oxygenation; HIT: heparin-induced thrombocytopenia.

The emerging use of extensive extracorporeal life support (ECLS), e.g. bridge to transplantation, requires novel approaches towards patient anticoagulation. This chapter provides an overview of the current evidence on this topic.

Background

ECLS (venoarterial) and extracorporeal membrane oxygenation (ECMO; venovenous) systems are designed for the temporary support of the failing heart and/or lungs. ECLS systems are small, closed priming volume systems, including a centrifugal pump and coated tubing without any option for removal of air. A cell saver device replaces cardiotomy suction. ECLS and ECMO require long-term anticoagulation, which is associated with an increased risk of thrombosis or bleeding.²⁹¹ The majority of patients exposed to long-term ECC develop acquired von Willebrand syndrome.

UFH anticoagulation is the standard in ECLS/ECMO, with similar dosing challenges to those experienced during cardiac surgery.^{292,293} Reagents, clot detection systems and reference values of different aPTT and ACT assays used during ECLS/ECMO vary greatly.^{291–293} Clotting tests do not take into account disturbances of the coagulation system, which are especially frequent in patients with multiple-organ failure requiring ECLS/ECMO support. Strict heparin dosing based on anti-Xa levels may increase the risk of excessive anticoagulation and bleeding.^{294,295} During the use of AT-independent anticoagulants, such as argatroban and bivalirudin, excessive anticoagulation and bleeding are avoided. However, in the absence of antidotes, bivalirudin requires haemofiltration to be eliminated from the circulation.²⁹⁶ Moreover, there is no strategy for enhanced extracorporeal elimination of argatroban, which is predominantly eliminated via the hepatic system.²⁹⁷

Description of the Evidence

There are significant institutional variations in the anticoagulation management and monitoring of ECLS/ECMO.²⁹⁸ The dose of heparin anticoagulation depends on the assay, with broad target ranges for the aPTT of 45–80 s and ACT of

140–220 s. Bivalirudin (0.02–0.05 mg/kg/h) and argatroban (0.1–0.2 mg/kg/h) are usually dosed on the basis of the aPTT with target ranges of 45–80 s and 50–60 s, respectively.

Particularly in adult patients, data are limited to small, mostly retrospective, single-centre studies. Clinical outcome studies are rare and mostly too underpowered to allow one to draw valid conclusions. In 1 study with 22 patients on ECMO, low anti-Xa levels were associated with an increase in deep venous thrombosis.²⁹⁹ This study also showed that the correlation of ACT or aPTT values with anti-Xa levels was poor to moderate, respectively.²⁹⁹ In 71 patients, a reduction of the ACT target value from 180–220 to 140–160 s reduced bleeding complications and transfusions without increasing thromboembolic events.³⁰⁰ The role of AT replacement in ECLS/ECMO remains to be clarified, because the evidence is limited to paediatric and small studies.^{298,301} Implementation of viscoelastic tests for anticoagulation monitoring during ECLS/ECMO appears to be promising but needs further validation.^{298,302} Successful use of bivalirudin^{303,304} and argatroban³⁰⁵ during ECLS/ECMO therapy based on aPTT dosing has been reported in small series in patients with and without HIT. Bivalirudin anticoagulation, when compared to heparin, resulted in more stable conditions of the haemostatic system and reduced bleeding and transfusion rates of bivalirudin.^{303,304}

In summary, the evidence for an optimal anticoagulation strategy during ECLS/ECMO is weak, and larger prospective studies are required.

8. Final Remarks

PBM in cardiac surgery comprises a multidisciplinary and multifactorial approach and consists of a sequence of separate strategies and interventions. A key factor in a successful PBM programme is the close collaboration between surgeons, anaesthetists, clinical perfusionists and intensivists. The joint effort of EACTS and EACTA to write this guideline underlines the importance of team efforts, and future guidelines should be extended to enable collaborations with other societies.

Our goal was to provide evidence-based recommendations with respect to blood-conservation strategies, preoperative patient optimization, intraoperative interventions and post-operative monitoring and treatment of bleeding disorders. All recommendations were based on bleeding-related end points, which narrowed our search strategy and reference list (Fig. 4). The lack of consensus on the definition of bleeding outcomes and their relevance for patient recovery warrants a consensus definition for bleeding complications in cardiac surgery for future studies. We also aimed to adhere to recommendations provided by other guidelines in this area.

A closer evaluation of the recommendations included in this guideline suggests that several fundamental aspects of PBM programmes are supported by the moderate quality of the evidence, and large RCTs are warranted to improve the evidence base of this guideline.

Additionally, it has to be highlighted that a large number of studies included in this guideline, in particular haemovigilance

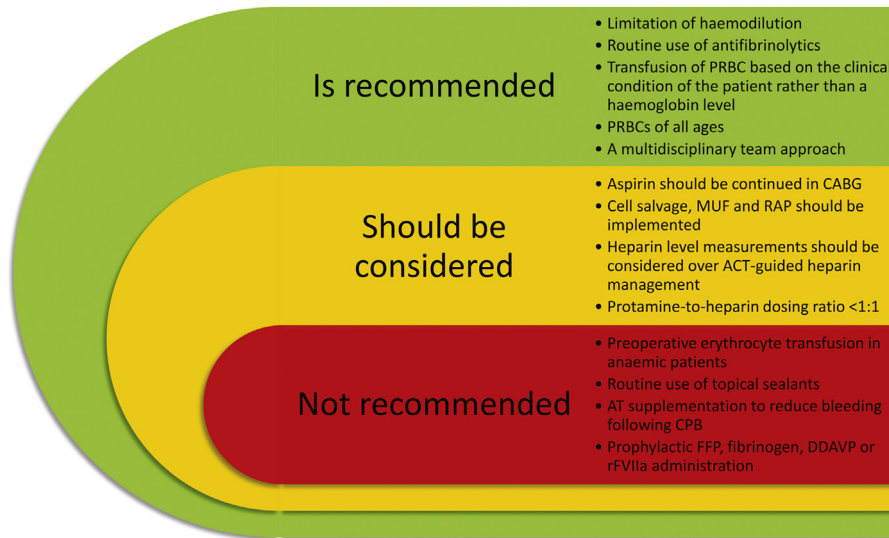


Figure 4. Key messages of the Joint Effort Patient Blood Management Guidelines for Adult Cardiac Surgery. ACT: activated clotting time; AT: antithrombin; CABG: coronary artery bypass grafting; CPB: cardiopulmonary bypass; DDAVP: desmopressin; FFP: fresh-frozen plasma; MUF: modified ultrafiltration; PRBC: packed red blood cells; RAP: retrograde autologous priming; rFVIIa: recombinant factor VIIa.

data, are derived from non-European countries. National regulations for the processing and leucodepletion of PRBCs differ among countries, and it is therefore unclear whether non-European data can be translated to the setting of European patients. We hope that the current guidelines represent a further step towards the establishment of a European standard for the quality of blood products, a European haemovigilance database and European standards for PBM.

Conflict of interest: 2014–2017: Umberto Benedetto, Daniel Bolliger, Andreas Koster, Michael Meesters, Domenico Pagano, Hanne Berg Ravn, Alexander Wahba, Milan Milojevic and Ruben Osnabrügge have nothing to disclose. Christa Boer attended consultancy meetings (Nordic Pharma) and received speaker's honoraria from LivaNova Group and Medtronic, Inc. Christian von Heymann received honoraria for lectures or consultancy work and travel reimbursements unrelated to his topic in this guideline from Bayer AG, Boehringer Ingelheim, Pfizer GmbH, Bristol Myers Squibb, Daiichi Sankyo, CSL Behring, Shire, Novo Nordisk, Ferring, Nordic Pharma, Mylan Healthcare GmbH, Sanofi-Aventis and HICC GbR during the last 3 years. Anders Jeppsson was a consultant for AstraZeneca, Boehringer-Ingelheim, XVIVO Perfusion, and LFB Corporation, and received speaker honoraria from AstraZeneca, CSL Behring, Roche Diagnostics, Triolab AB and Octapharma. Marco Ranucci was a consultant for LivaNova Group and received speaker's honoraria from Medtronic, Roche Diagnostics, TEM International, Vifor Pharma, Edwards Life Sciences, CSL Behring and Grifols. Alexander B.A. Vonk attended consultancy meetings (Nordic Pharma).

References

- Karkouti K, Wijeyesundera DN, Yau TM, Beattie WS, Abdelnaem E, McCluskey SA, et al. The independent association of massive blood loss with mortality in cardiac surgery. *Transfusion* 2004;44:1453–62.
- Ranucci M, Bozzetti G, Ditta A, Cotza M, Carboni G, Ballotta A. Surgical reexploration after cardiac operations: why a worse outcome? *Ann Thorac Surg* 2008;86:1557–62.
- Paone G, Likosky DS, Brewer R, Theurer PF, Bell GF, Cogan CM, et al. Transfusion of 1 and 2 units of red blood cells is associated with increased morbidity and mortality. *Ann Thorac Surg* 2014;97:87–93; discussion 93–4.
- Frojd V, Jeppsson A. Reexploration for bleeding and its association with mortality after cardiac surgery. *Ann Thorac Surg* 2016;102:109–17.
- Ndrepepa G, Schuster T, Hadamitzky M, Byrne RA, Mehilli J, Neumann FJ, et al. Validation of the Bleeding Academic Research Consortium definition of bleeding in patients with coronary artery disease undergoing percutaneous coronary intervention. *Circulation* 2012;125:1424–31.
- Mehran R, Rao SV, Bhatt DL, Gibson CM, Caixeta A, Eikelboom J, et al. Standardized bleeding definitions for cardiovascular clinical trials. A Consensus Report from the Bleeding Academic Research Consortium. *Circulation* 2011;123:2736–47.
- Alghamdi AA, Davis A, Brister S, Corey P, Logan A. Development and validation of Transfusion Risk Understanding Scoring Tool (TRUST) to stratify cardiac surgery patients according to their blood transfusion needs. *Transfusion* 2006;46:1120–9.
- Ranucci M, Aronson S, Dietrich W, Dyke CM, Hofmann A, Karkouti K, et al. Patient blood management during cardiac surgery: do we have enough evidence for clinical practice? *J Thorac Cardiovasc Surg* 2011;142:249.e1–32.
- Ferraris VA, Brown JR, Despotis GJ, Hammon JW, Reece TB, Saha SP, et al. 2011 update to the Society of Thoracic Surgeons and the Society of Cardiovascular Anesthesiologists blood conservation clinical practice guidelines. *Ann Thorac Surg* 2011;91:944–82.
- Goudie R, Sterne JAC, Verheyden V, Bhabra M, Ranucci M, Murphy GJ, et al. Risk scores to facilitate preoperative prediction of transfusion and large volume blood transfusion associated with adult cardiac surgery. *Br J Anaesth* 2015;114:757–66.
- Vuylsteke A, Pagel C, Gerrard C, Reddy B, Nashef S, Aldam P, et al. The Papworth Bleeding Risk Score: a stratification scheme for identifying cardiac surgery patients at risk of excessive early postoperative bleeding. *Eur J Cardiothorac Surg* 2011;39:924–30.
- Karkouti K, O'Farrell R, Yau TM, Beattie WS; Reducing Bleeding in Cardiac Surgery Research Group. Prediction of massive blood transfusion in cardiac surgery. *Can J Anesth* 2006;53:781–94.
- Al-Khabori M, Al-Riyami AZ, Mukaddirov M, Al-Sabti H. Transfusion indication predictive score: a proposed risk stratification score for

- perioperative red blood cell transfusion in cardiac surgery. *Vox Sang* 2014;107:269–75.
- 14 Biancari F, Brascia D, Onorati F, Reichart D, Perrotti A, Ruggieri VG, et al. Prediction of severe bleeding after coronary surgery: the WILL-BLEED Risk Score. *Thromb Haemost* 2017;117:445–56.
 - 15 Ranucci M, Castelveccchio S, Frigiola A, Scolletta S, Giomarelli P, Biagioli. Predicting transfusions in cardiac surgery: the easier, the better: the Transfusion Risk and Clinical Knowledge score. *Vox Sang* 2009;96:324–32.
 - 16 Kozek-Langenecker SA, Ahmed AB, Afshari A, Albaladejo P, Aldecoa C, Barauskas G, et al. Management of severe perioperative bleeding: guidelines from the European Society of Anaesthesiology: first update 2016. *Eur J Anaesthesiol* 2017;34:332–95.
 - 17 American Society of Anesthesiologists Task Force on Perioperative Blood Management. Practice guidelines for perioperative blood management: an updated report by the American Society of Anesthesiologists Task Force on Perioperative Blood Management. *Anesthesiology* 2015;122:241–75.
 - 18 Sousa-Uva M, Head SJ, Thielmann M, Cardillo G, Benedetto U, Czerny M, et al. Methodology manual for European Association for Cardio Thoracic Surgery (EACTS) clinical guidelines. *Eur J Cardiothorac Surg* 2015;48:809–16.
 - 19 Cantrill JA, Sibbald B, Buetow S. The Delphi and nominal group techniques in health services research. *Int J Pharm Pract* 1996;4:67–74.
 - 20 Mahla E, Suarez TA, Bliden KP, Rehak P, Metzler H, Sequeira AJ, et al. Platelet function measurement-based strategy to reduce bleeding and waiting time in clopidogrel-treated patients undergoing coronary artery bypass graft surgery: the timing based on platelet function strategy to reduce clopidogrel-associated bleeding related to CABG (TARGET-CABG) study. *Circ Cardiovasc Interv* 2012;5:261–9.
 - 21 Malm CJ, Hansson EC, Akeson J, Andersson M, Hesse C, Shams Hakim C, et al. Preoperative platelet function predicts perioperative bleeding complications in ticagrelor-treated cardiac surgery patients: a prospective observational study. *Br J Anaesth* 2016;117:309–15.
 - 22 Mahla E, Pruehler F, Farzi S, Pregartner G, Raggam RB, Beran E, et al. Does platelet reactivity predict bleeding in patients needing urgent coronary artery bypass grafting during dual antiplatelet therapy?. *Ann Thorac Surg* 2016;102:2010–7.
 - 23 Blome M, Isgro F, Kiessling AH, Skuras J, Haubelt H, Hellstern P, et al. Relationship between factor XIII activity, fibrinogen, haemostasis screening tests and postoperative bleeding in cardiopulmonary bypass surgery. *Thromb Haemost* 2005;93:1101–7.
 - 24 Bosch Y, Al Dieri R, ten Cate H, Nelemans P, Bloemen S, Hemker C, et al. Preoperative thrombin generation is predictive for the risk of blood loss after cardiac surgery: a research article. *J Cardiothorac Surg* 2013;8:154.
 - 25 Karlsson M, Ternstrom L, Hyllner M, Baghaei F, Nilsson S, Jeppsson A. Plasma fibrinogen level, bleeding, and transfusion after on-pump coronary artery bypass grafting surgery: a prospective observational study. *Transfusion* 2008;48:2152–8.
 - 26 Ranucci M, Jeppsson A, Baryshnikova E. Pre-operative fibrinogen supplementation in cardiac surgery patients: an evaluation of different trigger values. *Acta Anaesthesiol Scand* 2015;59:427–33.
 - 27 Walden K, Jeppsson A, Nasic S, Backlund E, Karlsson M. Low preoperative fibrinogen plasma concentration is associated with excessive bleeding after cardiac operations. *Ann Thorac Surg* 2014;97:1199–206.
 - 28 Gielen C, Dekkers O, Stijnen T, Schoones J, Brand A, Klautz R, et al. The effects of pre- and postoperative fibrinogen levels on blood loss after cardiac surgery: a systematic review and meta-analysis. *Interact CardioVasc Thorac Surg* 2014;18:292–8.
 - 29 Coakley M, Hall JE, Evans C, Duff E, Billing V, Yang L, et al. Assessment of thrombin generation measured before and after cardiopulmonary by-pass surgery and its association with postoperative bleeding. *J Thromb Haemost* 2011;9:282–92.
 - 30 Reinhöfer M, Brauer M, Franke U, Barz D, Marx G, Lösche W. The value of rotation thromboelastometry to monitor disturbed perioperative haemostasis and bleeding risk in patients with cardiopulmonary bypass. *Blood Coagul Fibrinolysis* 2008;19:212–9.
 - 31 Petricevic M, Biocina B, Milicic D, Konosic S, Svetina L, Lekic A, et al. Bleeding risk assessment using whole blood impedance aggregometry and rotational thromboelastometry in patients following cardiac surgery. *J Thromb Thrombolysis* 2013;36:514–26.
 - 32 Ranucci M, Baryshnikova E, Soro G, Ballotta A, De Benedetti D, Conti D. Multiple electrode whole-blood aggregometry and bleeding in cardiac surgery patients receiving thienopyridines. *Ann Thorac Surg* 2011;91:123–9.
 - 33 Ranucci M, Colella D, Baryshnikova E, Di Dedda U, Hemmings HC. Surgical, Clinical Outcome Research Group. Effect of preoperative P2Y12 and thrombin platelet receptor inhibition on bleeding after cardiac surgery. *Br J Anaesth* 2014;113:970–6.
 - 34 Mangano DT, Multicenter Study of Perioperative Ischemia Research Group. Aspirin and mortality from coronary bypass surgery. *N Engl J Med* 2002;347:1309–17.
 - 35 Musleh G, Dunning J. Does aspirin 6 h after coronary artery bypass grafting optimise graft patency? *Interact CardioVasc Thorac Surg* 2003;2:413–5.
 - 36 Smith PK, Goodnough LT, Levy JH, Poston RS, Short MA, Weerakkody GJ, et al. Mortality benefit with prasugrel in the TRITON-TIMI 38 coronary artery bypass grafting cohort: risk-adjusted retrospective data analysis. *J Am Coll Cardiol* 2012;60:388–96.
 - 37 Hansson EC, Jideus L, Aberg B, Bjursten H, Dreifaldt M, Holmgren A, et al. Coronary artery bypass grafting-related bleeding complications in patients treated with ticagrelor or clopidogrel: a nationwide study. *Eur Heart J* 2016;37:189–97.
 - 38 Gherli R, Mariscalco G, Dalen M, Onorati F, Perrotti A, Chocron S, et al. Safety of preoperative use of ticagrelor with or without aspirin compared with aspirin alone in patients with acute coronary syndromes undergoing coronary artery bypass grafting. *JAMA Cardiol* 2016;1:921–8.
 - 39 Douketis JD, Spyropoulos AC, Kaatz S, Becker RC, Caprini JA, Dunn AS, et al. Perioperative bridging anticoagulation in patients with atrial fibrillation. *N Engl J Med* 2015;373:823–33.
 - 40 O'Donnell MJ, Kearon C, Johnson J, Robinson M, Zondag M, Turpie I, et al. Brief communication: preoperative anticoagulant activity after bridging low-molecular-weight heparin for temporary interruption of warfarin. *Ann Intern Med* 2007;146:184–7.
 - 41 Medalion B, Frenkel G, Patachenko P, Hauptman E, Sasson L, Schachner A. Preoperative use of enoxaparin is not a risk factor for postoperative bleeding after coronary artery bypass surgery. *J Thorac Cardiovasc Surg* 2003;126:1875–9.
 - 42 Landenhed M, Johansson M, Erlinge D, Olsson ML, Bjursten H. Fondaparinux or enoxaparin: a comparative study of postoperative bleeding in coronary artery bypass grafting surgery. *Scand Cardiovasc J* 2010;44:100–6.
 - 43 Spyropoulos AC, Turpie AG, Dunn AS, Spandorfer J, Douketis J, Jacobson A, et al. Clinical outcomes with unfractionated heparin or low-molecular-weight heparin as bridging therapy in patients on long-term oral anticoagulants: the REGIMEN registry. *J Thromb Haemost* 2006;4:1246–52.
 - 44 Jones HU, Muhlestein JB, Jones KW, Bair TL, Lavasani F, Sohrevardi M, et al. Preoperative use of enoxaparin compared with unfractionated heparin increases the incidence of re-exploration for postoperative bleeding after open-heart surgery in patients who present with an acute coronary syndrome: clinical investigation and reports. *Circulation* 2002;106(12 Suppl 1):I19–22.
 - 45 Spyropoulos AC, Turpie AG, Dunn AS, Kaatz S, Douketis J, Jacobson A, et al. Perioperative bridging therapy with unfractionated heparin or low-molecular-weight heparin in patients with mechanical prosthetic heart valves on long-term oral anticoagulants (from the REGIMEN Registry). *Am J Cardiol* 2008;102:883–9.
 - 46 Douketis JD, Johnson JA, Turpie AG. Low-molecular-weight heparin as bridging anticoagulation during interruption of warfarin: assessment of a standardized periprocedural anticoagulation regimen. *Arch Intern Med* 2004;164:1319–26.
 - 47 Antithrombotic Trialists Collaboration. Collaborative meta-analysis of randomised trials of antiplatelet therapy for prevention of death,

- myocardial infarction, and stroke in high risk patients. *Br Med J* 2002;324:71–86.
- 48 Hastings S, Myles P, McIlroy D. Aspirin and coronary artery surgery: a systematic review and meta-analysis. *Br J Anaesth* 2015;115:376–85.
 - 49 Sun JC, Whitlock R, Cheng J, Eikelboom JW, Thabane L, Crowther MA, et al. The effect of pre-operative aspirin on bleeding, transfusion, myocardial infarction, and mortality in coronary artery bypass surgery: a systematic review of randomized and observational studies. *Eur Heart J* 2008;29:1057–71.
 - 50 Myles PS, Smith JA, Forbes A, Silbert B, Jayarajah M, Painter T, et al. Stopping vs. continuing aspirin before coronary artery surgery. *N Engl J Med* 2016;374:728–37.
 - 51 Deja MA, Kargul T, Domaradzki W, Staćel T, Mazur W, Wojakowski W. Effects of preoperative aspirin in coronary artery bypass grafting: a double-blind, placebo-controlled, randomized trial. *J Thorac Cardiovasc Surg* 2012;144:204–9.
 - 52 Morawski W, Sanak M, Cisowski M, Szczeklik M, Szczeklik W, Dropinski J, et al. Prediction of the excessive perioperative bleeding in patients undergoing coronary artery bypass grafting: role of aspirin and platelet glycoprotein IIIa polymorphism. *J Thorac Cardiovasc Surg* 2005;130:791–6.
 - 53 Hansson EC, Shams Hakimi C, Astrom-Olsson K, Hesse C, Wallen H, Dellborg M, et al. Effects of ex vivo platelet supplementation on platelet aggregability in blood samples from patients treated with acetylsalicylic acid, clopidogrel, or ticagrelor. *Br J Anaesth* 2014;112:570–5.
 - 54 Martin AC, Berndt C, Calmette L, Philip I, Decouture B, Gaussem P, et al. The effectiveness of platelet supplementation for the reversal of ticagrelor-induced inhibition of platelet aggregation: an in-vitro study. *Eur J Anaesthesiol* 2016;33:361–7.
 - 55 Yusuf S, Zhao F, Mehta SR, Chrolavicius S, Tognoni G, Fox KK. Effects of clopidogrel in addition to aspirin in patients with acute coronary syndromes without ST-segment elevation. *N Engl J Med* 2001;345:494–502.
 - 56 Wallentin L, Becker RC, Budaj A, Cannon CP, Emanuelsson H, Held C, et al. Ticagrelor versus clopidogrel in patients with acute coronary syndromes. *N Engl J Med* 2009;361:1045–57.
 - 57 Wiviott SD, Braunwald E, McCabe CH, Montalescot G, Ruzyllo W, Gottlieb S, et al. Prasugrel versus clopidogrel in patients with acute coronary syndromes. *N Engl J Med* 2007;357:2001–15.
 - 58 Becker RC, Bassand JP, Budaj A, Wojdyla DM, James SK, Cornel JH, et al. Bleeding complications with the P2Y₁₂ receptor antagonists clopidogrel and ticagrelor in the PLATElet inhibition and patient Outcomes (PLATO) trial. *Eur Heart J* 2011;32:2933–44.
 - 59 Levine GN, Bates ER, Bittl JA, Brindis RG, Fihn SD, Fleisher LA, et al. 2016 ACC/AHA guideline focused update on duration of dual antiplatelet therapy in patients with coronary artery disease. A Report of the American College of Cardiology/American Heart Association Task Force on Clinical Practice Guidelines. *J Am Coll Cardiol* 2016;68:1082–115.
 - 60 Qamar A, Bhatt DL. Current status of data on cangrelor. *Pharmacol Ther* 2016;159:102–9.
 - 61 Fox KA, Mehta SR, Peters R, Zhao F, Lakkis N, Gersh BJ, et al. Benefits and risks of the combination of clopidogrel and aspirin in patients undergoing surgical revascularization for non-ST-elevation acute coronary syndrome: the Clopidogrel in Unstable angina to prevent Recurrent ischemic Events (CURE) Trial. *Circulation* 2004;110:1202–8.
 - 62 Held C, Asenblad N, Bassand JP, Becker RC, Cannon CP, Claeys MJ, et al. Ticagrelor versus clopidogrel in patients with acute coronary syndromes undergoing coronary artery bypass surgery: results from the PLATO (Platelet Inhibition and Patient Outcomes) trial. *J Am Coll Cardiol* 2011;57:672–84.
 - 63 Tomsic A, Schotborgh MA, Manshanden JS, Li WW, de Mol BA. Coronary artery bypass grafting-related bleeding complications in patients treated with dual antiplatelet treatment. *Eur J Cardiothorac Surg* 2016;50:849–56.
 - 64 Pickard AS, Becker RC, Schumock GT, Frye CB. Clopidogrel-associated bleeding and related complications in patients undergoing coronary artery bypass grafting. *Pharmacotherapy* 2008;28:376–92.
 - 65 Purkayastha S, Athanasiou T, Malinowski V, Tekkis P, Foale R, Casula R, et al. Does clopidogrel affect outcome after coronary artery bypass grafting? A meta-analysis. *Heart* 2006;92:531–2.
 - 66 Kolh P, Windecker S, Alfonso F, Collet JP, Cremer J, Falk V, et al. 2014 ESC/EACTS Guidelines on myocardial revascularization: the Task Force on Myocardial Revascularization of the European Society of Cardiology (ESC) and the European Association for Cardio-Thoracic Surgery (EACTS). Developed with the special contribution of the European Association of Percutaneous Cardiovascular Interventions (EAPCI). *Eur J Cardiothorac Surg* 2014;46:517–92.
 - 67 Ferraris VA, Saha SP, Oestreich JH, Song HK, Rosengart T, Reece TB, et al. 2012 update to the Society of Thoracic Surgeons guideline on use of antiplatelet drugs in patients having cardiac and noncardiac operations. *Ann Thorac Surg* 2012;94:1761–81.
 - 68 Wallentin L. P2Y₁₂ inhibitors: differences in properties and mechanisms of action and potential consequences for clinical use. *Eur Heart J* 2009;30:1964–77.
 - 69 Gurbel PA, Bliden KP, Butler K, Tantry US, Gesheff T, Wei C, et al. Randomized double-blind assessment of the ONSET and OFFSET of the antiplatelet effects of ticagrelor versus clopidogrel in patients with stable coronary artery disease: the ONSET/OFFSET study. *Circulation* 2009;120:2577–85.
 - 70 Storey RF, Bliden KP, Ecob R, Karunakaran A, Butler K, Wei C, et al. Earlier recovery of platelet function after discontinuation of treatment with ticagrelor compared with clopidogrel in patients with high antiplatelet responses. *J Thromb Haemost* 2011;9:1730–7.
 - 71 Kwak YL, Kim JC, Choi YS, Yoo KJ, Song Y, Shim JK. Clopidogrel responsiveness regardless of the discontinuation date predicts increased blood loss and transfusion requirement after off-pump coronary artery bypass graft surgery. *J Am Coll Cardiol* 2010;56:1994–2002.
 - 72 Savonitto S, D'Urbano M, Caracciolo M, Barlocco F, Mariani G, Nichelatti M, et al. Urgent surgery in patients with a recently implanted coronary drug-eluting stent: a phase II study of 'bridging' antiplatelet therapy with tirofiban during temporary withdrawal of clopidogrel. *Br J Anaesth* 2010;104:285–91.
 - 73 Patrono C, Collier B, FitzGerald GA, Hirsh J, Roth G. Platelet-active drugs: the relationships among dose, effectiveness, and side effects: the Seventh ACCP Conference on Antithrombotic and Thrombolytic Therapy. *Chest* 2004;126:234S–64S.
 - 74 Dyke CM, Jennings LK, Maier G, Andreou C, Daly R, Tamberella MR. Preoperative platelet inhibition with eptifibatide during coronary artery bypass grafting with cardiopulmonary bypass. *J Cardiovasc Pharmacol Ther* 2007;12:54–60.
 - 75 De Carlo M, Maselli D, Cortese B, Ciabatti N, Gistri R, Levantino M, et al. Emergency coronary artery bypass grafting in patients with acute myocardial infarction treated with glycoprotein IIb/IIIa receptor inhibitors. *Int J Cardiol* 2008;123:229–33.
 - 76 Boeken U, Litmathe J, Kurt M, Feindt P, Gams E. CABG-procedures in patients with pretreatment with the GPIIb/IIIa-receptor antagonist tirofiban (Aggrastat): modification of perioperative management? *Int J Cardiol* 2008;127:257–9.
 - 77 Kincaid EH, Monroe ML, Saliba DL, Kon ND, Byerly WG, Reichert MG. Effects of preoperative enoxaparin versus unfractionated heparin on bleeding indices in patients undergoing coronary artery bypass grafting. *Ann Thorac Surg* 2003;76:124–8; discussion 8.
 - 78 Ashikhmina E, Tomasello N, Connors JM, Jahanyar J, Davidson M, Mizuguchi KA. Type A aortic dissection in a patient on dabigatran: hemostasis post circulatory arrest. *Ann Thorac Surg* 2014;98:2215–6.
 - 79 Stein P, Bosshart M, Brand B, Schlicker A, Spahn DR, Bettex D. Dabigatran anticoagulation and Stanford type A aortic dissection: lethal coincidence: case report with literature review. *Acta Anaesthesiol Scand* 2014;58:630–7.
 - 80 Crapelli GB, Bianchi P, Isgro G, Biondi A, de Vincentiis C, Ranucci M. A case of fatal bleeding following emergency surgery on an ascending aorta intramural hematoma in a patient taking dabigatran. *J Cardiothorac Vasc Anesth* 2016;30:1027–31.
 - 81 Weltert L, Rondinelli B, Bello R, Falco M, Bellisario A, Maselli D, et al. A single dose of erythropoietin reduces perioperative transfusions in cardiac surgery: results of a prospective single-blind randomized controlled trial. *Transfusion* 2015;55:1644–54.

- 82 Weltert L, D'Alessandro S, Nardella S, Girola F, Bellisario A, Maselli D, et al. Preoperative very short-term, high-dose erythropoietin administration diminishes blood transfusion rate in off-pump coronary artery bypass: a randomized blind controlled study. *J Thorac Cardiovasc Surg* 2010;139:621–6; discussion 6–7.
- 83 Yoo YC, Shim JK, Kim JC, Jo YY, Lee JH, Kwak YL. Effect of single recombinant human erythropoietin injection on transfusion requirements in preoperatively anemic patients undergoing valvular heart surgery. *Anesthesiology* 2011;115:929–37.
- 84 Hogan M, Klein AA, Richards T. The impact of anaemia and intravenous iron replacement therapy on outcomes in cardiac surgery. *Eur J Cardiothorac Surg* 2015;47:218–26.
- 85 Matsuda S, Fukui T, Shimizu J, Takao A, Takanashi S, Tomoike H. Associations between preoperative anemia and outcomes after off-pump coronary artery bypass grafting. *Ann Thorac Surg* 2013;95:854–60.
- 86 Zhang L, Hiebert B, Zarychanski R, Arora RC; Cardiovascular Health Research in Manitoba Investigator Group. Preoperative anemia does not increase the risks of early surgical revascularization after myocardial infarction. *Ann Thorac Surg* 2013;95:542–7.
- 87 Cutrell JB, Barros N, McBroom M, Luby J, Minhajuddin A, Ring WS, et al. Risk factors for deep sternal wound infection after cardiac surgery: influence of red blood cell transfusions and chronic infection. *Am J Infect Control* 2016;44:1302–9.
- 88 Engoren M, Schwann TA, Habib RH, Neill SN, Vance JL, Likosky DS. The independent effects of anemia and transfusion on mortality after coronary artery bypass. *Ann Thorac Surg* 2014;97:514–20.
- 89 von Heymann C, Kaufner L, Sander M, Spies C, Schmidt K, Gombotz H, et al. Does the severity of preoperative anemia or blood transfusion have a stronger impact on long-term survival after cardiac surgery? *J Thorac Cardiovasc Surg* 2016;152:1412–20.
- 90 Garrido-Martin P, Nassar-Mansur MI, de la Llana-Ducros R, Virgos-Aller TM, Rodriguez Fortuñez PM, Avalos-Pinto R, et al. The effect of intravenous and oral iron administration on perioperative anaemia and transfusion requirements in patients undergoing elective cardiac surgery: a randomized clinical trial. *Interact CardioVasc Thorac Surg* 2012;15:1013–8.
- 91 Johansson PI, Rasmussen AS, Thomsen LL. Intravenous iron isomaltoside 1000 (Monofer(R)) reduces postoperative anaemia in preoperatively non-anaemic patients undergoing elective or subacute coronary artery bypass graft, valve replacement or a combination thereof: a randomized double-blind placebo-controlled clinical trial (the PROTECT trial). *Vox Sang* 2015;109:257–66.
- 92 Kulier A, Levin J, Moser R, Rumpold-Seitlinger G, Tudor IC, Snyder-Ramos SA, et al. Impact of preoperative anemia on outcome in patients undergoing coronary artery bypass graft surgery. *Circulation* 2007;116:471–9.
- 93 Karkouti K, Wijeyesundera DN, Beattie WS; Reducing Bleeding in Cardiac Surgery Investigators. Risk associated with preoperative anemia in cardiac surgery: a multicenter cohort study. *Circulation* 2008;117:478–84.
- 94 Kozek-Langenecker SA, Afshari A, Albaladejo P, Santullano CA, De Robertis E, Filipesco DC, et al. Management of severe perioperative bleeding: guidelines from the European Society of Anaesthesiology. *Eur J Anaesthesiol* 2013;30:270–382.
- 95 Menkis AH, Martin J, Cheng DC, Fitzgerald DC, Freedman JJ, Gao C, et al. Drug, devices, technologies, and techniques for blood management in minimally invasive and conventional cardiothoracic surgery: a consensus statement from the International Society for Minimally Invasive Cardiothoracic Surgery (ISMICS) 2011. *Innovations (Phila)* 2012;7:229–41.
- 96 Karkouti K, Wijeyesundera DN, Yau TM, McCluskey SA, Chan CT, Wong PY, et al. Advance targeted transfusion in anemic cardiac surgical patients for kidney protection: an unblinded randomized pilot clinical trial. *Anesthesiology* 2012;116:613–21.
- 97 Deppe AC, Arbash W, Kuhn EW, Slottoch I, Scherner M, Liakopoulos OJ, et al. Current evidence of coronary artery bypass grafting off-pump versus on-pump: a systematic review with meta-analysis of over 16, 900 patients investigated in randomized controlled trials. *Eur J Cardiothorac Surg* 2016;49:1031–41; discussion 41.
- 98 Puskas JD, Martin J, Cheng DC, Benussi S, Bonatti JO, Diegeler A, et al. ISMICS consensus conference and statements of randomized controlled trials of off-pump versus conventional coronary artery bypass surgery. *Innovations (Phila)* 2015;10:219–29.
- 99 Anastasiadis K, Antonitsis P, Haidich AB, Argiriadou H, Deliopoulos A, Papakostantinou C. Use of minimal extracorporeal circulation improves outcome after heart surgery: a systematic review and meta-analysis of randomized controlled trials. *Int J Cardiol* 2013;164:158–69.
- 100 Harling L, Warren OJ, Martin A, Kemp PR, Evans PC, Darzi A, et al. Do miniaturized extracorporeal circuits confer significant clinical benefit without compromising safety? A meta-analysis of randomized controlled trials. *ASAIO J* 2011;57:141–51.
- 101 Sundermann SH, Sromicki J, Rodriguez Cetina Bieffer H, Seifert B, Holubec T, Falk V, et al. Mitral valve surgery: right lateral minithoracotomy or sternotomy? A systematic review and meta-analysis. *J Thorac Cardiovasc Surg* 2014;148:1989–95. e4.
- 102 Falk V, Cheng DC, Martin J, Diegeler A, Folliquet TA, Nifong LW, et al. Minimally invasive versus open mitral valve surgery: a consensus statement of the international society of minimally invasive coronary surgery (ISMICS) 2010. *Innovations (Phila)* 2011;6:66–76.
- 103 Phan K, Xie A, Di Eusanio M, Yan TD. A meta-analysis of minimally invasive versus conventional sternotomy for aortic valve replacement. *Ann Thorac Surg* 2014;98:1499–511.
- 104 Tavilla G, Bruggemans EF, Gielen CL, Brand A, van den Hout WB, Klautz RJ, et al. Multicentre randomized clinical trial to investigate the costeffectiveness of an allogeneic single-donor fibrin sealant after coronary artery bypass grafting (FIBER Study). *Br J Surg* 2015;102:1338–47.
- 105 Lamy A, Devereaux PJ, Prabhakaran D, Taggart DP, Hu S, Paolasso E, et al. Off-pump or on-pump coronary-artery bypass grafting at 30 days. *N Engl J Med* 2012;366:1489–97.
- 106 Diegeler A, Borgermann J, Kappert U, Breuer M, Boning A, Ursulescu A, et al. Off-pump versus on-pump coronary-artery bypass grafting in elderly patients. *N Engl J Med* 2013;368:1189–98.
- 107 Anastasiadis K, Murkin J, Antonitsis P, Bauer A, Ranucci M, Gygax E, et al. Use of minimal invasive extracorporeal circulation in cardiac surgery: principles, definitions and potential benefits. A position paper from the Minimal Invasive Extra-Corporeal Technologies international Society (MiECTiS). *Interact CardioVasc Thorac Surg* 2016;22:647–62.
- 108 Carless PA, Henry DA, Anthony DM. Fibrin sealant use for minimizing peri-operative allogeneic blood transfusion. *Cochrane Database Syst Rev* 2003;2:CD004171.
- 109 Abrishami A, Chung F, Wong J. Topical application of antifibrinolytic drugs for on-pump cardiac surgery: a systematic review and meta-analysis. *Can J Anaesth* 2009;56:202–12.
- 110 Ker K, Beecher D, Roberts I. Topical application of tranexamic acid for the reduction of bleeding. *Cochrane Database Syst Rev* 2013;7:D010562.
- 111 Gurian DB, Meneghini A, Abreu LC, Murad N, Matos LL, Pires AC, et al. A randomized trial of the topical effect of antifibrinolytic epsilon aminocaproic Acid on coronary artery bypass surgery without cardiopulmonary bypass. *Clin Appl Thromb Hemost* 2014;20:615–20.
- 112 Casalino S, Stelian E, Novelli E, De Jong A, Renzi L, Arellano F, et al. Reduced transfusion requirements with a closed cardiopulmonary bypass system. *J Cardiovasc Surg (Torino)* 2008;49:363–9.
- 113 Nakahira A, Sasaki Y, Hirai H, Matsuo M, Morisaki A, Suehiro S, et al. Cardiotomy suction, but not open venous reservoirs, activates coagulofibrinolysis in coronary artery surgery. *J Thorac Cardiovasc Surg* 2011;141:1289–97.
- 114 Ranucci M, Balduini A, Ditta A, Boncilli A, Brozzi S. A systematic review of biocompatible cardiopulmonary bypass circuits and clinical outcome. *Ann Thorac Surg* 2009;87:1311–9.
- 115 Landis RC, Brown JR, Fitzgerald D, Likosky DS, Shore-Lesserson L, Baker RA, et al. Attenuating the systemic inflammatory response to adult cardiopulmonary bypass: a critical review of the evidence base. *J Extra Corpor Technol* 2014;46:197–211.
- 116 Mahmood S, Bilal H, Zaman M, Tang A. Is a fully heparin-bonded cardiopulmonary bypass circuit superior to a standard cardiopulmonary bypass circuit? *Interact CardioVasc Thorac Surg* 2012;14:406–14.

- 117 Carless PA, Henry DA, Moxey AJ, O'Connell D, Brown T, Fergusson DA. Cell salvage for minimising perioperative allogeneic blood transfusion. *Cochrane Database Syst Rev* 2010;4:CD001888.
- 118 Vermeijden WJ, Hagenaars JA, Scheeren TW, de Vries AJ. Additional postoperative cell salvage of shed mediastinal blood in cardiac surgery does not reduce allogeneic blood transfusions: a cohort study. *Perfusion* 2016;31:384–90.
- 119 Vonk AB, Meesters MI, Garnier RP, Romijn JW, van Barneveld LJ, Heymans MW, et al. Intraoperative cell salvage is associated with reduced postoperative blood loss and transfusion requirements in cardiac surgery: a cohort study. *Transfusion* 2013;53:2782–9.
- 120 Zakkar M, Guida G, Angelini GD. Modified ultrafiltration in adult patients undergoing cardiac surgery. *Interact CardioVasc Thorac Surg* 2015;20:415–21.
- 121 Boodhwani M, Williams K, Babaev A, Gill G, Saleem N, Rubens FD. Ultrafiltration reduces blood transfusions following cardiac surgery: a meta-analysis. *Eur J Cardiothorac Surg* 2006;30:892–7.
- 122 Luciani GB, Menon T, Vecchi B, Auriemma S, Mazzucco A. Modified ultrafiltration reduces morbidity after adult cardiac operations: a prospective, randomized clinical trial. *Circulation* 2001;104:I253–9.
- 123 Sun P, Ji B, Sun Y, Zhu X, Liu J, Long C, et al. Effects of retrograde autologous priming on blood transfusion and clinical outcomes in adults: a meta-analysis. *Perfusion* 2013;28:238–43.
- 124 Saczkowski R, Bernier PL, Tchervenkov CI, Arellano R. Retrograde autologous priming and allogeneic blood transfusions: a meta-analysis. *Interact CardioVasc Thorac Surg* 2009;8:373–6.
- 125 Vandewiele K, Bove T, De Somer FM, Dujardin D, Vanackere M, De Smet D, et al. The effect of retrograde autologous priming volume on haemodilution and transfusion requirements during cardiac surgery. *Interact CardioVasc Thorac Surg* 2013;16:778–83.
- 126 Ranucci M, Baryshnikova E, Simeone F, Ranucci M, Scolletta S. Moderate-degree acidosis is an independent determinant of postoperative bleeding in cardiac surgery. *Minerva Anestesiol* 2015;81:885–93.
- 127 Lee BR, Song JW, Kwak YL, Yoo KJ, Shim JK. The influence of hypothermia on transfusion requirement in patients who received clopidogrel in proximity to off-pump coronary bypass surgery. *Yonsei Med J* 2014;55:224–31.
- 128 El-Sabbagh AM, Toomasian CJ, Toomasian JM, Ulyse G, Major T, Bartlett RH. Effect of air exposure and suction on blood cell activation and hemolysis in an in vitro cardiomy suction model. *ASAIO J* 2013;59:474–9.
- 129 Murphy GS, Hessel 2nd EA, Groom RC. Optimal perfusion during cardiopulmonary bypass: an evidence-based approach. *Anesth Analg* 2009;108:1394–417.
- 130 Lindholm L, Westerberg M, Bengtsson A, Ekroth R, Jensen E, Jeppsson A. A closed perfusion system with heparin coating and centrifugal pump improves cardiopulmonary bypass biocompatibility in elderly patients. *Ann Thorac Surg* 2004;78:2131–8; discussion 8.
- 131 Mirow N, Zittermann A, Koertke H, Maleszka A, Knobl H, Coskun T, et al. Heparin-coated extracorporeal circulation in combination with low dose systemic heparinization reduces early postoperative blood loss in cardiac surgery. *J Cardiovasc Surg (Torino)* 2008;49:277–84.
- 132 Paparella D, Scarscia G, Rotunno C, Marraudino N, Guida P, De Palo M, et al. A biocompatible cardiopulmonary bypass strategy to reduce hemostatic and inflammatory alterations: a randomized controlled trial. *J Cardiothorac Vasc Anesth* 2012;26:557–62.
- 133 Lorusso R, De Cicco G, Totaro P, Gelsomino S. Effects of phosphorylcholine coating on extracorporeal circulation management and postoperative outcome: a double-blind randomized study. *Interact CardioVasc Thorac Surg* 2008;8:7–11.
- 134 Marguerite S, Levy F, Quessard A, Dupeyron JP, Gros C, Steib A. Impact of a phosphorylcholine-coated cardiac bypass circuit on blood loss and platelet function: a prospective, randomized study. *J Extra Corpor Technol* 2012;44:5–9.
- 135 Hosoyama K, Ito K, Kawamoto S, Kumagai K, Akiyama M, Adachi O, et al. Poly-2-methoxyethylacrylate-coated cardiopulmonary bypass circuit can reduce transfusion of platelet products compared to heparincoated circuit during aortic arch surgery. *J Artif Organs* 2016;19:233–40.
- 136 Thiara AS, Mollnes TE, Videm V, Andersen VY, Svennevig K, Kolset SO, et al. Biocompatibility and pathways of initial complement pathway activation with Phisioand PMEA-coated cardiopulmonary bypass circuits during open-heart surgery. *Perfusion* 2011;26:107–14.
- 137 Carless PA, Henry DA, Moxey AJ, O'Connell DL, Brown T, Fergusson DA. Cell salvage for minimising perioperative allogeneic blood transfusion. *Cochrane Database Syst Rev* 2006;4:CD001888.
- 138 Wang G, Bainbridge D, Martin J, Cheng D. The efficacy of an intraoperative cell saver during cardiac surgery: a meta-analysis of randomized trials. *Anesth Analg* 2009;109:320–30.
- 139 Weltert L, Nardella S, Rondinelli MB, Pierelli L, De Paulis R. Reduction of allogeneic red blood cell usage during cardiac surgery by an integrated intraand postoperative blood salvage strategy: results of a randomized comparison. *Transfusion* 2013;53:790–7.
- 140 Engels GE, van Klarenbosch J, Gu YJ, van Oeveren W, de Vries AJ. Intraoperative cell salvage during cardiac surgery is associated with reduced postoperative lung injury. *Interact CardioVasc Thorac Surg* 2016;22:298–304.
- 141 Gabel J, Westerberg M, Bengtsson A, Jeppsson A. Cell salvage of cardiomy suction blood improves the balance between proand anti-inflammatory cytokines after cardiac surgery. *Eur J Cardiothorac Surg* 2013;44:506–11.
- 142 Damgaard S, Nielsen CH, Andersen LW, Bendtzen K, Tvede M, Steinbruchel DA. Cell saver for on-pump coronary operations reduces systemic inflammatory markers: a randomized trial. *Ann Thorac Surg* 2010;89:1511–7.
- 143 Jewell AE, Akowuah EF, Suvarna SK, Braidley P, Hopkinson D, Cooper G. A prospective randomised comparison of cardiomy suction and cell saver for recycling shed blood during cardiac surgery. *Eur J Cardiothorac Surg* 2003;23:633–6.
- 144 Rubens FD, Boodhwani M, Mesana T, Wozny D, Wells G, Nathan HJ. The cardiomy trial: a randomized, double-blind study to assess the effect of processing of shed blood during cardiopulmonary bypass on transfusion and neurocognitive function. *Circulation* 2007;116:189–97.
- 145 Al-Khabori M, Al-Riyami AZ, Baskaran B, Siddiqi M, Al-Sabti H. Discriminatory power of the intraoperative cell salvage use in the prediction of platelet and plasma transfusion in patients undergoing cardiac surgery. *Transfus Apher Sci* 2015;53:208–12.
- 146 Padhi S, Kemmis-Betty S, Rajesh S, Hill J, Murphy MF, Guideline Development Group. Blood transfusion: summary of NICE guidance. *Br Med J* 2015;351:h5832.
- 147 Liumbruno GM, Waters JH. Unwashed shed blood: should we transfuse it?. *Blood Transfus* 2011;9:241–5.
- 148 Hou X, Yang F, Liu R, Yang J, Zhao Y, Wan C, et al. Retrograde autologous priming of the cardiopulmonary bypass circuit reduces blood transfusion in small adults: a prospective, randomized trial. *Eur J Anaesthesiol* 2009;26:1061–6.
- 149 Nathan HJ, Parlea L, Dupuis JY, Hendry P, Williams KA, Rubens FD, et al. Safety of deliberate intraoperative and postoperative hypothermia for patients undergoing coronary artery surgery: a randomized trial. *J Thorac Cardiovasc Surg* 2004;127:1270–5.
- 150 Vonk AB, Veerhoek D, van den Brom CE, van Barneveld LJ, Boer C. Individualized heparin and protamine management improves rotational thromboelastometric parameters and postoperative hemostasis in valve surgery. *J Cardiothorac Vasc Anesth* 2014;28:235–41.
- 151 Noui N, Zogheib E, Walczak K, Werbrout A, Amar AB, Dupont H, et al. Anticoagulation monitoring during extracorporeal circulation with the Hepcon/HMS device. *Perfusion* 2012;27:214–20.
- 152 Hoenicka M, Rupp P, Muller-Eising K, Deininger S, Kunert A, Liebold A, et al. Anticoagulation management during multivessel coronary artery bypass grafting: a randomized trial comparing individualized heparin management and conventional hemostasis management. *J Thromb Haemost* 2015;13:1196–206.
- 153 Wang J, Ma HP, Zheng H. Blood loss after cardiopulmonary bypass, standard vs titrated protamine: a meta-analysis. *Neth J Med* 2013;71:123–7.
- 154 Meesters MI, Veerhoek D, de Lange F, de Vries JW, de Jong JR, Romijn JW, et al. Effect of high or low protamine dosing on postoperative

- bleeding following heparin anticoagulation in cardiac surgery. A randomised clinical trial. *Thromb Haemost* 2016;116:251–61.
- 155 Avidan MS, Levy JH, Scholz J, Delphin E, Rosseel PM, Howie MB, et al. A phase III, double-blind, placebo-controlled, multicenter study on the efficacy of recombinant human antithrombin in heparin-resistant patients scheduled to undergo cardiac surgery necessitating cardiopulmonary bypass. *Anesthesiology* 2005;102:276–84.
 - 156 Avidan MS, Levy JH, van Aken H, Feneck RO, Latimer RD, Ott E, et al. Recombinant human antithrombin III restores heparin responsiveness and decreases activation of coagulation in heparin-resistant patients during cardiopulmonary bypass. *J Thorac Cardiovasc Surg* 2005;130:107–13.
 - 157 Finley A, Greenberg C. Review article: heparin sensitivity and resistance: management during cardiopulmonary bypass. *Anesth Analg* 2013;116:1210–22.
 - 158 Hofmann B, Bushnaq H, Kraus FB, Raspe C, Simm A, Silber RE, et al. Immediate effects of individualized heparin and protamine management on hemostatic activation and platelet function in adult patients undergoing cardiac surgery with tranexamic acid antifibrinolytic therapy. *Perfusion* 2013;28:412–8.
 - 159 Aziz KA, Masood O, Hoschitzky JA, Ronald A. Does use of the Hepcon point-of-care coagulation monitor to optimise heparin and protamine dosage for cardiopulmonary bypass decrease bleeding and blood and blood product requirements in adult patients undergoing cardiac surgery? *Interact Cardio Vasc Thorac Surg* 2006;5:469–82.
 - 160 Pappalardo F, Franco A, Crescenzi G, De Simone F, Torracca L, Zangrillo A. Anticoagulation management in patients undergoing open heart surgery by activated clotting time and whole blood heparin concentration. *Perfusion* 2006;21:285–90.
 - 161 Avidan MS, Alcock EL, Da Fonseca J, Ponte J, Desai JB, Despotis GJ, et al. Comparison of structured use of routine laboratory tests or near-patient assessment with clinical judgement in the management of bleeding after cardiac surgery. *Br J Anaesth* 2004;92:178–86.
 - 162 Slight RD, Buell R, Nzewi OC, McClelland DB, Mankad PS. A comparison of activated coagulation time-based techniques for anticoagulation during cardiac surgery with cardiopulmonary bypass. *J Cardiothorac Vasc Anesth* 2008;22:47–52.
 - 163 Radulovic V, Laffin A, Hansson KM, Backlund E, Baghaei F, Jeppsson A. Heparin and protamine titration does not improve haemostasis after cardiac surgery: a prospective randomized study. *PLoS One* 2015;10:e0130271.
 - 164 Teoh KH, Young E, Blackall MH, Roberts RS, Hirsh J. Can extra protamine eliminate heparin rebound following cardiopulmonary bypass surgery? *J Thorac Cardiovasc Surg* 2004;128:211–9.
 - 165 Koster A, Borgermann J, Gummert J, Rudloff M, Zittermann A, Schirmer U. Protamine overdose and its impact on coagulation, bleeding, and transfusions after cardiopulmonary bypass: results of a randomized double-blind controlled pilot study. *Clin Appl Thromb Hemost* 2014;20:290–5.
 - 166 Slaughter TF, Mark JB, El-Moalem H, Hayward KA, Hilton AK, Hodgins LP, et al. Hemostatic effects of antithrombin III supplementation during cardiac surgery: results of a prospective randomized investigation. *Blood Coagul Fibrinolysis* 2001;12:25–31.
 - 167 Ranucci M, Baryshnikova E, Crapelli GB, Woodward MK, Paez A, Pelissero G. Preoperative antithrombin supplementation in cardiac surgery: a randomized controlled trial. *J Thorac Cardiovasc Surg* 2013;145:1393–9.
 - 168 Linkins LA, Dans AL, Moores LK, Bona R, Davidson BL, Schulman S, et al. Treatment and prevention of heparin-induced thrombocytopenia: antithrombotic therapy and prevention of thrombosis, 9th ed: American College of Chest Physicians Evidence-Based Clinical Practice Guidelines. *Chest* 2012;141:e495S–530.
 - 169 Warkentin TE, Sheppard JA. Serological investigation of patients with a previous history of heparin-induced thrombocytopenia who are reexposed to heparin. *Blood* 2014;123:2485–93.
 - 170 Lo GK, Sigouin CS, Warkentin TE. What is the potential for overdiagnosis of heparin-induced thrombocytopenia? *Am J Hematol* 2007;82:1037–43.
 - 171 Smedira NG, Dyke CM, Koster A, Jurmann M, Bhatia DS, Hu T, et al. Anticoagulation with bivalirudin for off-pump coronary artery bypass grafting: the results of the EVOLUTION-OFF study. *J Thorac Cardiovasc Surg* 2006;131:686–92.
 - 172 Dyke CM, Aldea G, Koster A, Smedira N, Avery E, Aronson S, et al. Off-pump coronary artery bypass with bivalirudin for patients with heparin-induced thrombocytopenia or antiplatelet factor four/heparin antibodies. *Ann Thorac Surg* 2007;84:836–9.
 - 173 Dyke CM, Smedira NG, Koster A, Aronson S, McCarthy 2nd HL, Kirshner R, et al. A comparison of bivalirudin to heparin with protamine reversal in patients undergoing cardiac surgery with cardiopulmonary bypass: the EVOLUTION-ON study. *J Thorac Cardiovasc Surg* 2006;131:533–9.
 - 174 Koster A, Dyke CM, Aldea G, Smedira NG, McCarthy 2nd HL, Aronson S, et al. Bivalirudin during cardiopulmonary bypass in patients with previous or acute heparin-induced thrombocytopenia and heparin antibodies: results of the CHOOSE-ON trial. *Ann Thorac Surg* 2007;83:572–7.
 - 175 Warkentin TE, Greinacher A, Koster A. Bivalirudin. *Thromb Haemost* 2008;99:830–9.
 - 176 Palatianos G, Michalis A, Alivizatos P, Lacoumenda S, Geroulanos S, Karabinis A, et al. Perioperative use of iloprost in cardiac surgery patients diagnosed with heparin-induced thrombocytopenia-reactive antibodies or with true HIT (HIT-reactive antibodies plus thrombocytopenia): an 11-year experience. *Am J Hematol* 2015;90:608–17.
 - 177 Koster A, Meyer O, Fischer T, Kukucka M, Krabatsch T, Bauer M, et al. One-year experience with the platelet glycoprotein IIb/IIIa antagonist tirofiban and heparin during cardiopulmonary bypass in patients with heparin-induced thrombocytopenia type II. *J Thorac Cardiovasc Surg* 2001;122:1254–5.
 - 178 Welsby IJ, Um J, Milano CA, Ortel TL, Arepally G. Plasmapheresis and heparin reexposure as a management strategy for cardiac surgical patients with heparin-induced thrombocytopenia. *Anesth Analg* 2010;110:30–5.
 - 179 Warkentin TE, Sheppard JA, Chu FV, Kapoor A, Crowther MA, Gangji A. Plasma exchange to remove HIT antibodies: dissociation between enzyme-immunoassay and platelet activation test reactivities. *Blood* 2015;125:195–8.
 - 180 Tran MH, Lin DM, Wilcox T, Schiro D, Cannesson M, Milliken J. Effects of a multimodality blood conservation schema toward improvement of intraoperative hemoglobin levels and off-pump transfusions in coronary artery bypass graft surgery. *Transfusion* 2014;54:2769–74.
 - 181 Vretzakis G, Kleitsaki A, Stamoulis K, Bareka M, Georgopoulou S, Karanikolas M, et al. Intra-operative intravenous fluid restriction reduces perioperative red blood cell transfusion in elective cardiac surgery, especially in transfusion-prone patients: a prospective, randomized controlled trial. *J Cardiothorac Surg* 2010;5:7.
 - 182 Ranucci M, Conti D, Castelvechio S, Menicanti L, Frigiola A, Ballotta A, et al. Hematocrit on cardiopulmonary bypass and outcome after coronary surgery in nontransfused patients. *Ann Thorac Surg* 2010;89:11–7.
 - 183 Martin K, Keller E, Gertler R, Tassani P, Wiesner G. Efficiency and safety of preoperative autologous blood donation in cardiac surgery: a matched-pair analysis in 432 patients. *Eur J Cardiothorac Surg* 2010;37:1396–401.
 - 184 Dietrich W, Thuermer K, Heyde S, Busley R, Berger K. Autologous blood donation in cardiac surgery: reduction of allogeneic blood transfusion and cost-effectiveness. *J Cardiothorac Vasc Anesth* 2005;19:589–96.
 - 185 Carless P, Moxey A, O'Connell D, Henry D. Autologous transfusion techniques: a systematic review of their efficacy. *Transfus Med* 2004;14:123–44.
 - 186 Barile L, Fominskiy E, Di Tomasso N, Alpizar Castro LE, Landoni G, De Luca M, et al. Acute normovolemic hemodilution reduces allogeneic red blood cell transfusion in cardiac surgery: a systematic review and metaanalysis of randomized trials. *Anesth Analg* 2017;124:743–52.
 - 187 Osawa EA, Rhodes A, Landoni G, Galas FR, Fukushima JT, Park CH, et al. Effect of perioperative goal-directed hemodynamic resuscitation

- therapy on outcomes following cardiac surgery: a randomized clinical trial and systematic review. *Crit Care Med* 2016;44:724–33.
- 188 Damen T, Reinsfelt B, Redfors B, Nygren A. Pressure-dependent changes in haematocrit and plasma volume during anaesthesia, a randomised clinical trial. *Acta Anaesthesiol Scand* 2016;60:560–8.
 - 189 Goepfert MS, Richter HP, Zu Eulenburg C, Gruetzmacher J, Rafflenbeul E, Roeher K, et al. Individually optimized hemodynamic therapy reduces complications and length of stay in the intensive care unit: a prospective, randomized controlled trial. *Anesthesiology* 2013;119:824–36.
 - 190 Bethlehem I, Wierda K, Visser C, Jekel L, Koopmans M, Kuiper MA. Influence of two colloidal extracorporeal primes on coagulation of cardiac surgical patients: a prospectively randomized open-label pilot trial. *J Extra Corpor Technol* 2014;46:293–9.
 - 191 Cho J-E, Shim J-K, Song J-W, Song J-W, Lee H-W, Kim D-H, et al. Effect of 6% hydroxyethyl starch 130/0.4 as a priming solution on coagulation and inflammation following complex heart surgery. *Yonsei Med J* 2014;55:625–34.
 - 192 Kuitunen AH, Hynynen MJ, Vahtera E, Salmenpera MT. Hydroxyethyl starch as a priming solution for cardiopulmonary bypass impairs hemostasis after cardiac surgery. *Anesth Analg* 2004;98:291–7.
 - 193 Schramko A, Suojaranta-Ylinen R, Niemi T, Pesonen E, Kuitunen A, Raivio P, et al. The use of balanced HES 130/0.42 during complex cardiac surgery; effect on blood coagulation and fluid balance: a randomized controlled trial. *Perfusion* 2015;30:224–32.
 - 194 Kamra C, Beney A. Human albumin in extracorporeal prime: effect on platelet function and bleeding. *Perfusion* 2013;28:536–40.
 - 195 Van der Linden PJ, De Hert SG, Daper A, Trenchant A, Schmartz D, Defrance P, et al. 3.5% urea-linked gelatin is as effective as 6% HES 200/0.5 for volume management in cardiac surgery patients. *Can J Anesth* 2004;51:236–41.
 - 196 Kasper SM, Meinert P, Kampe S, Gorg C, Geisen C, Mehlhorn U, et al. Large-dose hydroxyethyl starch 130/0.4 does not increase blood loss and transfusion requirements in coronary artery bypass surgery compared with hydroxyethyl starch 200/0.5 at recommended doses. *Anesthesiology* 2003;99:42–7.
 - 197 Gurbuz HA, Durukan AB, Salman N, Tavlasoglu M, Durukan E, Ucar HI, et al. Hydroxyethyl starch 6%, 130/0.4 vs. a balanced crystalloid solution in cardiopulmonary bypass priming: a randomized, prospective study. *J Cardiothorac Surg* 2013;8:71.
 - 198 Min JOS, Ramzisham ARM, Zamrin MD. Is 6% hydroxyethyl starch 130/0.4 safe in coronary artery bypass graft surgery? *Asian Cardiovasc Thorac Ann* 2009;17:368–72.
 - 199 Vanhoonacker J, Ongenaes M, Vanoverschelde H, Donadoni R. Hydroxyethyl starch 130/0.4 versus modified fluid gelatin for cardiopulmonary bypass priming: the effects on postoperative bleeding and volume expansion needs after elective CABG. *Acta Anaesthesiol Belg* 2009;60:91–7.
 - 200 Hecht-Dolnik M, Barkan H, Taharka A, Loftus J. Hetastarch increases the risk of bleeding complications in patients after off-pump coronary bypass surgery: a randomized clinical trial. *J Thorac Cardiovasc Surg* 2009;138:703–11.
 - 201 Van der Linden PJ, De Hert SG, Deraedt D, Cromheecke S, De Decker K, De Paep R, et al. Hydroxyethyl starch 130/0.4 versus modified fluid gelatin for volume expansion in cardiac surgery patients: the effects on perioperative bleeding and transfusion needs. *Anesth Analg* 2005;101:629–34.
 - 202 Skhirtladze K, Base EM, Lassnigg A, Kaider A, Linke S, Dworschak M, et al. Comparison of the effects of albumin 5%, hydroxyethyl starch 130/0.4 6%, and Ringer's lactate on blood loss and coagulation after cardiac surgery. *Br J Anaesth* 2014;112:255–64.
 - 203 DeFoe GR, Ross CS, Olmstead EM, Surgenor SD, Fillinger MP, Groom RC, et al. Lowest hematocrit on bypass and adverse outcomes associated with coronary artery bypass grafting. Northern New England Cardiovascular Disease Study Group. *Ann Thorac Surg* 2001;71:769–76.
 - 204 Lewis CE, Hiratzka LF, Woods SE, Hendy MP, Engel AM. Autologous blood transfusion in elective cardiac valve operations. *J Card Surg* 2005;20:513–8.
 - 205 Myles PS, Smith JA, Forbes A, Silbert B, Jayarajah M, Painter T, et al. Tranexamic acid in patients undergoing coronary-artery surgery. *N Engl J Med* 2017;376:136–48.
 - 206 Henry DA, Carless PA, Moxey AJ, O'Connell D, Stokes BJ, Fergusson DA, et al. Anti-fibrinolytic use for minimising perioperative allogeneic blood transfusion. *Cochrane Database Syst Rev* 2011;3:CD001886.
 - 207 Sedrakyan A, Treasure T, Eleftheriades JA. Effect of aprotinin on clinical outcomes in coronary artery bypass graft surgery: a systematic review and meta-analysis of randomized clinical trials. *J Thorac Cardiovasc Surg* 2004;128:442–8.
 - 208 Desborough M, Sandu R, Brunskill SJ, Doree C, Trivella M, Montedori A, et al. Fresh frozen plasma for cardiovascular surgery. *Cochrane Database Syst Rev* 2015;7:CD007614.
 - 209 Stanworth SJ, Brunskill SJ, Hyde CJ, McClelland DB, Murphy MF. Is fresh frozen plasma clinically effective? A systematic review of randomized controlled trials. *Br J Haematol* 2004;126:139–52.
 - 210 Casbard AC, Williamson LM, Murphy MF, Rege K, Johnson T. The role of prophylactic fresh frozen plasma in decreasing blood loss and correcting coagulopathy in cardiac surgery. A systematic review. *Anaesthesia* 2004;59:550–8.
 - 211 Demeyere R, Gillardin S, Arnout J, Strengers PF. Comparison of fresh frozen plasma and prothrombin complex concentrate for the reversal of oral anticoagulants in patients undergoing cardiopulmonary bypass surgery: a randomized study. *Vox Sang* 2010;99:251–60.
 - 212 Johansen M, Wikkelsø A, Lunde J, Wetterslev J, Afshari A. Prothrombin complex concentrate for reversal of vitamin K antagonist treatment in bleeding and non-bleeding patients. *Cochrane Database Syst Rev* 2015;7:CD010555.
 - 213 Godje O, Gallmeier U, Schelian M, Grunewald M, Mair H. Coagulation factor XIII reduces postoperative bleeding after coronary surgery with extracorporeal circulation. *Thorac Cardiovasc Surg* 2006;54:26–33.
 - 214 Karkouti K, von Heymann C, Jespersen CM, Korte W, Levy JH, Ranucci M, et al. Efficacy and safety of recombinant factor XIII on reducing blood transfusions in cardiac surgery: a randomized, placebo-controlled, multicenter clinical trial. *J Thorac Cardiovasc Surg* 2013;146:927–39.
 - 215 Bilecen S, de Groot JA, Kalkman CJ, Spanjersberg AJ, Brandon Bravo Bruinsma GJ, Moons KG, et al. Effect of fibrinogen concentrate on intraoperative blood loss among patients with intraoperative bleeding during high-risk cardiac surgery: a randomized clinical trial. *JAMA* 2017;317:738–47.
 - 216 Wikkelsø A, Lunde J, Johansen M, Stensballe J, Wetterslev J, Møller AM, et al. Fibrinogen concentrate in bleeding patients. *Cochrane Database Syst Rev* 2013;8:CD008864.
 - 217 Fassl J, Matt P, Eckstein F, Filipovic M, Gregor M, Zenklusen U, et al. Transfusion of allogeneic blood products in proximal aortic surgery with hypothermic circulatory arrest: effect of thromboelastometry-guided transfusion management. *J Cardiothorac Vasc Anesth* 2013;27:1181–8.
 - 218 Bilecen S, Peelen LM, Kalkman CJ, Spanjersberg AJ, Moons KG, Nierich AP. Fibrinogen concentrate therapy in complex cardiac surgery. *J Cardiothorac Vasc Anesth* 2013;27:12–7.
 - 219 Ortmann E, Besser MW, Sharples LD, Gerrard C, Berman M, Jenkins DP, et al. An exploratory cohort study comparing prothrombin complex concentrate and fresh frozen plasma for the treatment of coagulopathy after complex cardiac surgery. *Anesth Analg* 2015;121:26–33.
 - 220 Bolliger D, Gorlinger K, Tanaka KA. Pathophysiology and treatment of coagulopathy in massive hemorrhage and hemodilution. *Anesthesiology* 2010;113:1205–19.
 - 221 Bignami E, Cattaneo M, Crescenzi G, Ranucci M, Guarracino F, Cariello C, et al. Desmopressin after cardiac surgery in bleeding patients. A multicenter randomized trial. *Acta Anaesthesiol Scand* 2016;60:892–900.
 - 222 Wademan BH, Galvin SD. Desmopressin for reducing postoperative blood loss and transfusion requirements following cardiac surgery in adults. *Interact CardioVasc Thorac Surg* 2014;18:360–70.
 - 223 Gill R, Herbertson M, Vuylsteke A, Olsen PS, von Heymann C, Mythen M, et al. Safety and efficacy of recombinant activated factor VII: a randomized placebo-controlled trial in the setting of bleeding after cardiac surgery. *Circulation* 2009;120:21–7.
 - 224 Simpson E, Lin Y, Stanworth S, Birchall J, Doree C, Hyde C. Recombinant factor VIIa for the prevention and treatment of bleeding in patients without haemophilia. *Cochrane Database Syst Rev* 2012;3:CD005011.

- 225 Adler Ma SC, Brindle W, Burton G, Gallacher S, Hong FC, Manelius I, et al. Tranexamic acid is associated with less blood transfusion in offpump coronary artery bypass graft surgery: a systematic review and meta-analysis. *J Cardiothorac Vasc Anesth* 2011;25:26–35.
- 226 Ker K, Edwards P, Perel P, Shakur H, Roberts I. Effect of tranexamic acid on surgical bleeding: systematic review and cumulative meta-analysis. *Br Med J* 2012;344:e3054.
- 227 Sharma V, Katznelson R, Jerath A, Garrido-Olivares L, Carroll J, Rao V, et al. The association between tranexamic acid and convulsive seizures after cardiac surgery: a multivariate analysis in 11 529 patients. *Anaesthesia* 2014;69:124–30.
- 228 Koster A, Borgermann J, Zittermann A, Lueth JU, Gillis-Januszewski T, Schirmer U. Moderate dosage of tranexamic acid during cardiac surgery with cardiopulmonary bypass and convulsive seizures: incidence and clinical outcome. *Br J Anaesth* 2013;110:34–40.
- 229 Karkouti K, Beattie WS, Dattilo KM, McCluskey SA, Ghannam M, Hamdy A, et al. A propensity score case-control comparison of aprotinin and tranexamic acid in high-transfusion-risk cardiac surgery. *Transfusion* 2006;46:327–38.
- 230 Mangano DT, Tudor IC, Dietzel C, Multicenter Study of Perioperative Ischemia Research Group; Ischemia Research and Education Foundation. The risk associated with aprotinin in cardiac surgery. *N Engl J Med* 2006;354:353–65.
- 231 Walkden GJ, Verheyden V, Goudie R, Murphy GJ. Increased perioperative mortality following aprotinin withdrawal: a real-world analysis of blood management strategies in adult cardiac surgery. *Intensive Care Med* 2013;39:1808–17.
- 232 Howell N, Senanayake E, Freemantle N, Pagano D. Putting the record straight on aprotinin as safe and effective: results from a mixed treatment meta-analysis of trials of aprotinin. *J Thorac Cardiovasc Surg* 2013;145:234–40.
- 233 Pagano D, Howell NJ, Freemantle N, Cunningham D, Bonser RS, Graham TR, et al. Bleeding in cardiac surgery: the use of aprotinin does not affect survival. *J Thorac Cardiovasc Surg* 2008;135:495–502.
- 234 Fergusson DA, Hebert PC, Mazer CD, Fremes S, MacAdams C, Murkin JM, et al. A comparison of aprotinin and lysine analogues in high-risk cardiac surgery. *N Engl J Med* 2008;358:2319–31.
- 235 McMullan V, Alston RP. III. Aprotinin and cardiac surgery: a sorry tale of evidence misused. *Br J Anaesth* 2013;110:675–8.
- 236 Canada H. Health Canada Decision on Trasylol (Aprotinin) (21 September 2011, date last accessed).
- 237 Agency E. European Medicines Agency Recommends Lifting Suspension of Aprotinin; Review Finds That Benefits of All Antifibrinolytic Medicines Outweigh Risks in Restricted Range of Indications (17 February 2012, date last accessed).
- 238 Yang L, Stanworth S, Hopewell S, Doree C, Murphy M. Is fresh-frozen plasma clinically effective? An update of a systematic review of randomized controlled trials. *Transfusion* 2012;52:1673–86; quiz.
- 239 Karkouti K, McCluskey SA, Syed S, Pazaratz C, Poonawala H, Crowther MA. The influence of perioperative coagulation status on postoperative blood loss in complex cardiac surgery: a prospective observational study. *Anesth Analg* 2010;110:1533–40.
- 240 Ternstrom L, Radulovic V, Karlsson M, Baghaei F, Hyllner M, Bylock A, et al. Plasma activity of individual coagulation factors, hemodilution and blood loss after cardiac surgery: a prospective observational study. *Thromb Res* 2010;126:e128–33.
- 241 Theusinger OM, Baulig W, Asmis LM, Seifert B, Spahn DR. In vitro factor XIII supplementation increases clot firmness in Rotation Thromboelastometry (ROTEM). *Thromb Haemost* 2010;104:385–91.
- 242 Karkouti K, Callum J, Crowther MA, McCluskey SA, Pendergrast J, Tait G, et al. The relationship between fibrinogen levels after cardiopulmonary bypass and large volume red cell transfusion in cardiac surgery: an observational study. *Anesth Analg* 2013;117:14–22.
- 243 Levy JH, Gill R, Nussmeier NA, Olsen PS, Andersen HF, Booth FV, et al. Repletion of factor XIII following cardiopulmonary bypass using a recombinant A-subunit homodimer. A preliminary report. *Thromb Haemost* 2009;102:765–71.
- 244 Hiippala ST, Myllyla GJ, Vahtera EM. Hemostatic factors and replacement of major blood loss with plasma-poor red cell concentrates. *Anesth Analg* 1995;81:360–5.
- 245 Bolliger D, Gonsahn M, Levy JH, Williams WH, Tanaka KA. Is preoperative fibrinogen predictive for postoperative bleeding after coronary artery bypass grafting surgery? *Transfusion* 2009;49:2006–7; author reply 7–8.
- 246 Tanaka KA, Esper S, Bolliger D. Perioperative factor concentrate therapy. *Br J Anaesth* 2013;111(Suppl 1):i35–49.
- 247 Jeppsson A, Walden K, Roman-Emanuel C, Thimour-Bergstrom L, Karlsson M. Preoperative supplementation with fibrinogen concentrate in cardiac surgery: a randomized controlled study. *Br J Anaesth* 2016;116:208–14.
- 248 Rahe-Meyer N, Solomon C, Hanke A, Schmidt DS, Knoerzer D, Hochleitner G, et al. Effects of fibrinogen concentrate as first-line therapy during major aortic replacement surgery: a randomized, placebo-controlled trial. *Anesthesiology* 2013;118:40–50.
- 249 Ranucci M, Baryshnikova E, Crapelli GB, Rahe-Meyer N, Menicanti L, Frigiola A. Randomized, double-blinded, placebo-controlled trial of fibrinogen concentrate supplementation after complex cardiac surgery. *J Am Heart Assoc* 2015;4. e002066.
- 250 Rahe-Meyer N, Levy JH, Mazer CD, Schramko A, Klein AA, Brat R, et al. Randomized evaluation of fibrinogen vs placebo in complex cardiovascular surgery (REPLACE): a double-blind phase III study of haemostatic therapy. *Br J Anaesth* 2016;117:41–51.
- 251 Ranucci M, Baryshnikova E. Fibrinogen supplementation after cardiac surgery: insights from the Zero-Plasma trial (ZEPLAST). *Br J Anaesth* 2016;116:618–23.
- 252 Fassl J, Lurati Buse G, Filipovic M, Reuthebuch O, Hampf K, Seeberger MD, et al. Perioperative administration of fibrinogen does not increase adverse cardiac and thromboembolic events after cardiac surgery. *Br J Anaesth* 2015;114:225–34.
- 253 Ghadimi K, Levy JH, Welsby IJ. Prothrombin complex concentrates for bleeding in the perioperative setting. *Anesth Analg* 2016;122:1287–300.
- 254 Arnekian V, Camous J, Fattal S, Rezaiguia-Delclaux S, Nottin R, Stephan F. Use of prothrombin complex concentrate for excessive bleeding after cardiac surgery. *Interact CardioVasc Thorac Surg* 2012;15:382–9.
- 255 Cappabianca G, Mariscalco G, Biancari F, Maselli D, Papeso F, Cottini M, et al. Safety and efficacy of prothrombin complex concentrate as first-line treatment in bleeding after cardiac surgery. *Crit Care* 2016;20:5.
- 256 Carless PA, Henry DA, Moxey AJ, O'Connell D, McClelland B, Henderson KM, et al. Desmopressin for minimising perioperative allogeneic blood transfusion. *Cochrane Database Syst Rev* 2004;1:CD001884.
- 257 Crescenzi G, Landoni G, Biondi-Zoccai G, Pappalardo F, Nuzzi M, Bignami E, et al. Desmopressin reduces transfusion needs after surgery: a meta-analysis of randomized clinical trials. *Anesthesiology* 2008;109:1063–76.
- 258 Desborough MJ, Oakland KA, Landoni G, Crivellari M, Doree C, Estcourt LJ, et al. Desmopressin for treatment of platelet dysfunction and reversal of antiplatelet agents: a systematic review and meta-analysis of randomized controlled trials. *J Thromb Haemost* 2017;15:263–72.
- 259 Steiner ME, Ness PM, Assmann SF, Triulzi DJ, Sloan SR, Delaney M, et al. Effects of red-cell storage duration on patients undergoing cardiac surgery. *N Engl J Med* 2015;372:1419–29.
- 260 Lacroix J, Hebert PC, Fergusson DA, Tinmouth A, Cook DJ, Marshall JC, et al. Age of transfused blood in critically ill adults. *N Engl J Med* 2015;372:1410–8.
- 261 Dzieciatkowska M, Silliman CC, Moore EE, Kelher MR, Banerjee A, Land KJ, et al. Proteomic analysis of the supernatant of red blood cell units: the effects of storage and leucoreduction. *Vox Sang* 2013;105:210–8.
- 262 Membe SK, Coyle D, Husereau D, Cimon K, Tinmouth A, Normandin S. Octaplas Compared with Fresh Frozen Plasma to Reduce the Risk of Transmitting Lipid-Enveloped Viruses: An Economic Analysis and Budget Impact Analysis [Internet]. 2011.
- 263 Deppe AC, Weber C, Zimmermann J, Kuhn EW, Slottosch I, Liakopoulos OJ, et al. Point-of-care thromboelastography/thromboelastometry-based coagulation management in cardiac surgery: a meta-analysis of 8332 patients. *J Surg Res* 2016;203:424–33.

- 264 Serraino GF, Murphy GJ. Routine use of viscoelastic blood tests for diagnosis and treatment of coagulopathic bleeding in cardiac surgery: updated systematic review and meta-analysis. *Br J Anaesth* 2017;118:823–33.
- 265 Karkouti K, Callum J, Wijeyesundera DN, Rao V, Crowther M, Grocott HP, et al. Point-of-care hemostatic testing in cardiac surgery: a stepped-wedge clustered randomized controlled trial. *Circulation* 2016;134:1152–62.
- 266 Murphy GJ, Pike K, Rogers CA, Wordsworth S, Stokes EA, Angelini GD, et al. Liberal or restrictive transfusion after cardiac surgery. *N Engl J Med* 2015;372:997–1008.
- 267 Hajjar LA, Vincent JL, Galas FR, Nakamura RE, Silva CM, Santos MH, et al. Transfusion requirements after cardiac surgery: the TRACS randomized controlled trial. *JAMA* 2010;304:1559–67.
- 268 von Heymann C, Sander M, Foer A, Heinemann A, Spiess B, Braun J, et al. The impact of an hematocrit of 20% during normothermic cardiopulmonary bypass for elective low risk coronary artery bypass graft surgery on oxygen delivery and clinical outcome—a randomized controlled study [ISRCTN35655335]. *Crit Care* 2006;10:R58.
- 269 Klausen SS, Hervig T, Seghatchian J, Reikvam H. Bacterial contamination of blood components: Norwegian strategies in identifying donors with higher risk of inducing septic transfusion reactions in recipients. *Transfus Apher Sci* 2014;51:97–102.
- 270 Dwyre DM, Fernando LP, Holland PV. Hepatitis B, hepatitis C and HIV transfusion-transmitted infections in the 21st century. *Vox Sang* 2011;100:92–8.
- 271 Shaz BH, Stowell SR, Hillyer CD. Transfusion-related acute lung injury: from bedside to bench and back. *Blood* 2011;117:1463–71.
- 272 Dunbar NM. Current options for transfusion-related acute lung injury risk mitigation in platelet transfusions. *Curr Opin Hematol* 2015;22:554–8.
- 273 Vamvakas EC, Blajchman MA. Transfusion-related immunomodulation (TRIM): an update. *Blood Rev* 2007;21:327–48.
- 274 Blumberg N, Zhao H, Wang H, Messing S, Heal JM, Lyman GH. The intention-to-treat principle in clinical trials and meta-analyses of leukoreduced blood transfusions in surgical patients. *Transfusion* 2007;47: 573–81.
- 275 Rohde JM, Dimcheff DE, Blumberg N, Saint S, Langa KM, Kuhn L, et al. Health care-associated infection after red blood cell transfusion: a systematic review and meta-analysis. *JAMA* 2014;311:1317–26.
- 276 Bilgin YM, van de Watering LM, Eijssman L, Versteegh MI, Brand R, van Oers MH, et al. Double-blind, randomized controlled trial on the effect of leukocyte-depleted erythrocyte transfusions in cardiac valve surgery. *Circulation* 2004;109:2755–60.
- 277 Sartipy U, Holzmann MJ, Hjalgrim H, Edgren G. Red blood cell concentrate storage and survival after cardiac surgery. *JAMA* 2015;314: 1641–3.
- 278 Alexander PE, Barty R, Fei Y, Vandvik PO, Pai M, Siemieniuk RA, et al. Transfusion of fresher vs older red blood cells in hospitalized patients: a systematic review and meta-analysis. *Blood* 2016;127:400–10.
- 279 Vamvakas EC. Relative safety of pooled whole blood-derived versus single-donor (apheresis) platelets in the United States: a systematic review of disparate risks. *Transfusion* 2009;49:2743–58.
- 280 Welsby IJ, Lockhart E, Phillips-Bute B, Campbell ML, Mathew JP, Newman MF, et al. Storage age of transfused platelets and outcomes after cardiac surgery. *Transfusion* 2010;50:2311–7.
- 281 Ogawa S, Szlam F, Bolliger D, Nishimura T, Chen EP, Tanaka KA. The impact of hematocrit on fibrin clot formation assessed by rotational thromboelastometry. *Anesth Analg* 2012;115:16–21.
- 282 Wikkelso A, Wetterslev J, Moller AM, Afshari A. Thromboelastography (TEG) or thromboelastometry (ROTEM) to monitor haemostatic treatment versus usual care in adults or children with bleeding. *Cochrane Database Syst Rev* 2016;8:CD007871.
- 283 Whiting P, Al M, Westwood M, Ramos IC, Ryder S, Armstrong N, et al. Viscoelastic point-of-care testing to assist with the diagnosis, management and monitoring of haemostasis: a systematic review and costeffectiveness analysis. *Health Technol Assess* 2015;19:1–228. v–vi.
- 284 Weber CF, Gorlinger K, Meininger D, Herrmann E, Bingold T, Moritz A, et al. Point-of-care testing: a prospective, randomized clinical trial of efficacy in coagulopathic cardiac surgery patients. *Anesthesiology* 2012;117:531–47.
- 285 Song HK, von Heymann C, Jespersen CM, Karkouti K, Korte W, Levy JH, et al. Safe application of a restrictive transfusion protocol in moderate-risk patients undergoing cardiac operations. *Ann Thorac Surg* 2014;97:1630–5.
- 286 Ranucci M, Aloisio T, Carboni G, Ballotta A, Pistuddi V, Menicanti L, et al. Acute kidney injury and hemodilution during cardiopulmonary bypass: a changing scenario. *Ann Thorac Surg* 2015;100:95–100.
- 287 Patel NN, Avlonitis VS, Jones HE, Reeves BC, Sterne JA, Murphy GJ. Indications for red blood cell transfusion in cardiac surgery: a systematic review and meta-analysis. *Lancet Haematol* 2015;2:e543–53.
- 288 Nakamura RE, Vincent JL, Fukushima JT, de Almeida JP, Franco RA, Lee Park C, et al. A liberal strategy of red blood cell transfusion reduces cardiogenic shock in elderly patients undergoing cardiac surgery. *J Thorac Cardiovasc Surg* 2015;150:1314–20.
- 289 Kumar A, Mhaskar R, Grossman BJ, Kaufman RM, Tobian AA, Kleinman S, et al. Platelet transfusion: a systematic review of the clinical evidence. *Transfusion* 2015;55:1116–27. quiz 5.
- 290 Kaufman RM, Djulbegovic B, Gernsheimer T, Kleinman S, Tinmouth AT, Capocelli KE, et al. Platelet transfusion: a clinical practice guideline from the AABB. *Ann Intern Med* 2015;162:205–13.
- 291 Annich GM. Extracorporeal life support: the precarious balance of hemostasis. *J Thromb Haemost* 2015;13(Suppl 1):S336–42.
- 292 Bates SM, Weitz JI. Coagulation assays. *Circulation* 2005;112:e53–60.
- 293 Wool GD, Lu CM, Education Committee of the Academy of Clinical Laboratory Physicians and Scientists. Pathology consultation on anticoagulation monitoring: factor X-related assays. *Am J Clin Pathol* 2013;140:623–34.
- 294 Price EA, Jin J, Nguyen HM, Krishnan G, Bowen R, Zehnder JL. Discordant aPTT and anti-Xa values and outcomes in hospitalized patients treated with intravenous unfractionated heparin. *Ann Pharmacother* 2013;47:151–8.
- 295 Adatya S, Uriel N, Yarmohammadi H, Holley CT, Feng A, Roy SS, et al. Anti-factor Xa and activated partial thromboplastin time measurements for heparin monitoring in mechanical circulatory support. *JACC Heart Fail* 2015;3:314–22.
- 296 Koster A, Chew D, Grundel M, Hausmann H, Grauhan O, Kuppe H, et al. An assessment of different filter systems for extracorporeal elimination of bivalirudin: an in vitro study. *Anesth Analg* 2003;96:1316–9.
- 297 Hoechter DJ, von Dossow V, Winter H, Muller HH, Meiser B, Neurohr C, et al. The munich lung transplant group: intraoperative extracorporeal circulation in lung transplantation. *Thorac Cardiovasc Surg* 2015;63:706–14.
- 298 Bembea MM, Annich G, Rycus P, Oldenburg G, Berkowitz I, Pronovost P. Variability in anticoagulation management of patients on extracorporeal membrane oxygenation: an international survey. *Pediatr Crit Care Med* 2013;14:e77–84.
- 299 Chu DC, Abu-Samra AG, Baird GL, Devers C, Sweeney J, Levy MM, et al. Quantitative measurement of heparin in comparison with conventional anticoagulation monitoring and the risk of thrombotic events in adults on extracorporeal membrane oxygenation. *Intensive Care Med* 2015;41:369–70.
- 300 Yeo HJ, Kim do H, Jeon D, Kim YS, Cho WH. Low-dose heparin during extracorporeal membrane oxygenation treatment in adults. *Intensive Care Med* 2015;41:2020–1.
- 301 Byrnes JW, Swearingen CJ, Prodan P, Fiser R, Dyamenahalli U. Antithrombin III supplementation on extracorporeal membrane oxygenation: impact on heparin dose and circuit life. *ASAIO J* 2014;60:57–62.
- 302 Ranucci M, Baryshnikova E, Cotza M, Carboni G, Isgrò G, Carlucci C, et al. Coagulation monitoring in postcardiotomy ECMO: conventional tests, point-of-care, or both? *Minerva Anestesiol* 2016;82:858–66.
- 303 Ranucci M, Ballotta A, Kandil H, Isgrò G, Carlucci C, Baryshnikova E, et al. Bivalirudin-based versus conventional heparin anticoagulation for postcardiotomy extracorporeal membrane oxygenation. *Crit Care* 2011;15:R275.
- 304 Pieri M, Agracheva N, Bonaveglia E, Greco T, De Bonis M, Covello RD, et al. Bivalirudin versus heparin as an anticoagulant during extracorporeal membrane oxygenation: a case-control study. *J Cardiothorac Vasc Anesth* 2013;27:30–4.
- 305 Beiderlinden M, Treschan T, Gorlinger K, Peters J. Argatroban in extracorporeal membrane oxygenation. *Artif Organs* 2007;31:461–5.